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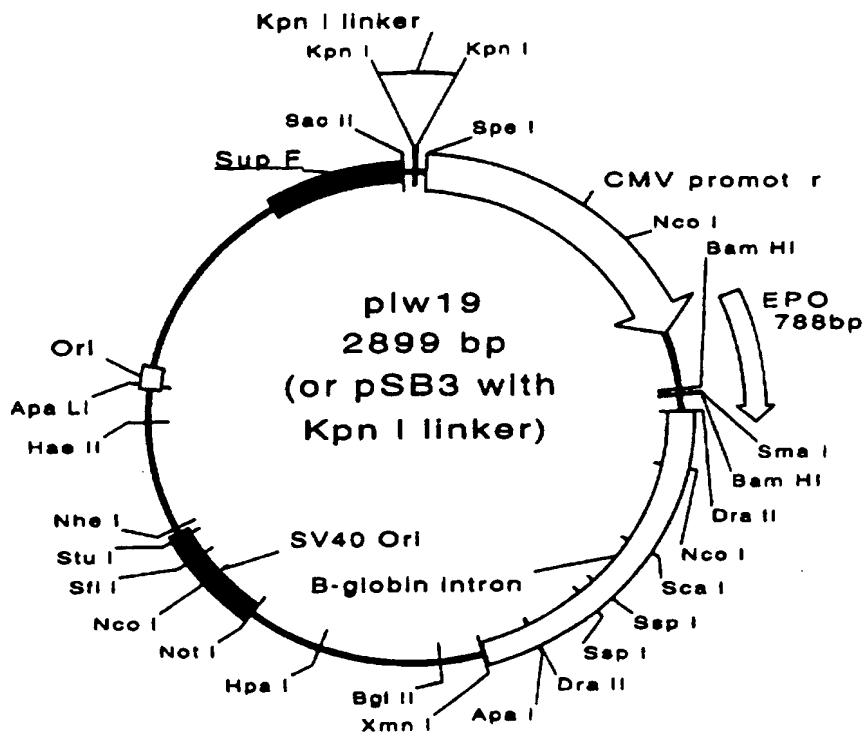
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(54) Title: RECOMBINANT DNA MOLECULES AND EXPRESSION VECTORS FOR ERYTHROPOIETIN

## (57) Abstract

A recombinant DNA molecule adapted for transfection of a host cell comprising a nucleic acid molecule encoding mammalian erythropoietin, an expression control sequence operatively linked thereto and at least one SAR element. The invention also relates to expression vectors having the recombinant DNA molecule and to mammalian cells transformed with the expression vector. The mammalian cells lack multiple copies of an amplified amplification gene and are capable of expressing recombinant EPO in vitro at levels of at least 1,500 u/10<sup>6</sup> cells in 24 hours. The invention further relates to a method of expressing recombinant mammalian erythropoietin using the expression vectors and to a transgenic non-human animal or embryo whose germ cells and somatic cells contain a DNA construct having the recombinant DNA molecule of the invention.



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**Title: Recombinant DNA Molecules and  
Expression Vectors for Erythropoietin**

**FIELD OF THE INVENTION**

The invention relates to recombinant DNA molecules  
5 adapted for transfection of a host cell, and having a  
nucleic acid molecule encoding mammalian erythropoietin,  
operatively linked to an expression control sequence and  
having at least one SAR element. The invention also  
relates to expression vectors for transfection of a host  
10 cell and to host cells for expressing erythropoietin.  
The invention further relates to methods of preparing  
recombinant erythropoietin using the host cells  
transfected with the expression vectors.

**BACKGROUND OF THE INVENTION**

15 Erythropoietin (EPO) is a heavily glycosylated  
acidic glycoprotein with a molecular weight of  
approximately 35,000. The protein consists of 166 amino  
acids and has a leader signal sequence of 27 amino acids  
which is removed *in vivo* during secretion from the host  
20 cell. The sequence encoding the unprocessed EPO is 579  
nucleotides in length (Jacobs et al, 1985, Lin et al,  
1985, U.S. Patent 4,703,008 and WO 86/03520).

Erythropoietin is the principal hormone involved in  
the regulation and maintenance of physiological levels of  
25 erythrocytes in mammalian circulation and functions to  
promote erythroid development, to initiate hemoglobin  
synthesis and to stimulate proliferation of immature  
erythroid precursors. The hormone is produced primarily  
by the adult kidney and foetal liver and is maintained in  
30 the circulation at concentrations of about 10-20  
milliunits/ml of serum under normal physiological  
conditions. Elevated levels of EPO, induced by tissue  
hypoxia, trigger proliferation and differentiation of a  
population of receptive progenitor stem cells in the bone  
35 marrow, stimulating hemoglobin synthesis in maturing

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erythroid cells and accelerating the release of erythrocytes from the marrow into the circulation.

Recombinant EPO has been used to successfully treat patients, including patients having anemia as a result of chronic renal failure. As EPO is the primary regulator of red blood cell formation, it has applications in both the diagnosis and treatment of disorders of red blood cell production and has potential applications for treating a range of conditions.

10 The urine of severely anaemic patients was, at one time, almost the sole source for the commercial isolation of EPO. U.S. Patent No. 3,033,753 describes a method for obtaining a crude EPO preparation from sheep plasma. The preparation of monoclonal antibodies specific for human EPO provided a means for identifying EPO produced from EPO mRNA, for screening libraries and for cloning the EPO gene. Human EPO cDNA has been cloned and expressed in E. coli (Lee-Huang, 1984, Proc. Natl. Acad. Sci. 81:2708). Isolation of the human EPO gene using mixtures of short or long synthetic nucleotides as probes led to the expression of biologically active EPO in mammalian cells (Lin, 1985, Proc. Natl. Acad. Sci. 82:7580; Lin, WO 85/02610; Jacobs, et al., 1985, Nature (Lond.) 313:806; Goto et al., 1988, Biotechnology 6:67). Jacobs, et al., 1985, supra, described the use of plasmids containing EPO DNA which were not integrated into the chromosomes of the COS host cells, but replicated autonomously in the cells to many thousands of copies, thereby killing the cells. Thus the expression of EPO was only a transient phenomenon in these cells.

Lin, in U.S. Patent No. 4,703,008, reported expression of the human EPO gene in COS-1 and CHO cells. However, attempts to use transfected cells as production vehicles for EPO have been hampered by the low levels of EPO expressed by transfected cells. Given the important applications of recombinant EPO, there is much interest in developing more efficient methods for the expression of

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EPO.

- Lin in U.S. Patent No. 4,703,008 reported methods to increase the low amounts of EPO produced by transfected CHO cells (e.g. 2.99u/ml/3 days) by a process of gene amplification. Levels of approximately 1500 units EPO/10<sup>6</sup> cells/48 hours were reported by Lin, following amplification. Gene amplification involves culturing cells in appropriate media conditions to select cells resistant to a selective agent, such as the drug methotrexate.
- Selection for cells resistant to methotrexate produces cells containing greater numbers of DHFR genes and passenger genes, such as the EPO gene carried on the expression vector along with the DHFR gene or transfected with the DHFR gene.
- However, gene amplification is a very time consuming and labour intensive process. A major disadvantage of amplification is the inherent instability of amplified genes (McDonald, 1990, Crit. Rev. Biotech. 10:155). As it is usually necessary to maintain the amplified cells in the presence of toxic analogs to maintain high copy number, amplification may be inappropriate for large scale production due to the costs and toxicity of the selective agent. The high copy number of the DHFR-target transgene may also sequester transcription factor, leading to a retardation of cell growth.

Genomic clones of human EPO have been used in attempts to develop stably transfected mammalian cell lines that secrete high levels of active erythropoietin (Powell, et al., 1986, Proc. Natl. Acad. Sci. 83:6465; Masatsuga, et al., European Patent Application Publication No. 0 236 059). In PCT Application WO 88/00241 Powell, describes the preparation of mammalian cell lines (COS-7 and BHK) transfected with the Apa I restriction fragment of the human EPO gene and selected for high expression by amplification.

Human EPO cDNA has also been expressed in mammalian cells (Yanagi, et al., 1989, DNA 8: 419). Bernstein, in

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PCT application WO 86/03520 describes the expression of EPO cDNA in various host cells, resulting in the secretion of up to 160 ng/ml of EPO into the medium after amplification. European Patent Application publication  
5 No. 0 267 678 discloses expression of recombinant EPO and secretion into the culture medium at levels of 600 units/ml.

A few scaffold attachment region (SAR) elements have been shown to increase the expression of reporter genes in  
10 transfected cells. SAR elements are thought to be DNA sequences which mediate attachment of chromatin loops to the nuclear matrix or scaffold. SAR elements are also known as MAR (matrix-associated regions) (reviewed by Phi-  
Van and Strätling, Prog. Mol. Subcell. Biol. 11:1-11,  
15 1990). These elements will hereinafter be referred to as "SAR elements". SAR elements are usually 300 or more base pairs long, and they require a redundancy of sequence information and contain multiple sites of protein-DNA interaction. SAR elements are found in non-coding  
20 regions: in flanking regions or introns.

Stief, et al., (Nature 341:343-345, 1989) stably transfected chicken macrophage cells by constructs which contained the CAT gene either fused to the lysozyme promoter, or to the lysozyme promoter and the lysozyme  
25 enhancer. When the transcription units contained in both constructs were flanked on both sides by lysozyme 5' SAR elements (A elements), gene expression was increased about 10 times relative to transfectants, which contained the constructs lacking the SAR elements.

30 Phi-Van, et al., (Mol. Cell. Biol., 10:2302-2307, 1990) determined the influence of the SAR element located 5' to the chicken lysozyme gene (A element) on the CAT gene expression from a heterologous promoter (herpes simplex virus thymidine kinase promoter) in stably  
35 transfected heterologous cells (rat fibroblasts). The median CAT activity per copy number in transfectants was 10 times higher for the transcriptional unit flanked on

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both sides by A elements than for the transcriptional unit lacking SAR elements.

Klehr, et al., (Biochemistry, 30:1264-1270, 1991) stably transfected mouse L cells by different constructs containing the human interferon  $\beta$  gene. When the construct was flanked by SAR elements, the gene transcription level was enhanced 20-30 fold with respect to the SAR-free construct, containing only the immediate regulatory elements.

However, the above-noted experiments have been limited to a very few examples of SAR elements, expressing mostly reporter genes, such as chloramphenicol acetyl transferase (CAT) or luciferase. SAR elements have not shown consistent results in their effect on the expression of target genes and some target gene sequences have been found to inhibit the effect of SAR elements (Klehr, et al., 1991, Biochemistry 30:1264).

#### SUMMARY OF THE INVENTION

The present inventor has significantly found that SAR elements may be used to increase the expression of recombinant mammalian EPO DNA. The present inventor constructed expression vectors carrying EPO genomic or cDNA sequences flanked by 3' and 5' human apolipoprotein B SAR elements. The expression vectors, when transfected into host cells resulted in increased expression of EPO compared to control host cells transfected with EPO expression vectors lacking SAR elements. Host cells transfected with expression vectors carrying an EPO cDNA sequence flanked by 3' and 5' SAR elements which expressed high levels of EPO were selected and cloned to obtain homogenous stable cell lines over-expressing EPO. Cloning produced stable cell lines expressing high levels of EPO, without the need for amplification. SAR elements have not, to the inventor's knowledge, heretofore been used for long term expression of a target gene in stable cell lines, or for the expression of recombinant EPO.

The present invention thus provides a recombinant

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DNA molecule adapted for transfection of a host cell comprising a nucleic acid molecule encoding mammalian erythropoietin, an expression control sequence operatively linked thereto and at least one SAR element. In an embodiment, the nucleic acid molecule encodes mammalian erythropoietin having the amino acid sequence shown in SEQ ID NOS 33 or 34 and Figure 3B. In another embodiment, the nucleic acid molecule has the sequence as shown in SEQ ID NO. 33 or 35.

10           The SAR element is preferably a SAR element co-mapping with the chromatin domain boundary, such as the human apolipoprotein SAR elements, most preferably the SAR element comprises the sequence as shown in SEQ ID NO. 36 or 37 and Figures 5 or 6.

15           The present invention also provides an expression vector comprising a recombinant DNA molecule adapted for transfection of a host cell comprising a nucleic acid molecule encoding mammalian erythropoietin, an expression control sequence operatively linked thereto and at least  
20 one SAR element.

In an embodiment, the expression vector comprises a nucleic acid molecule encoding erythropoietin and having the sequence shown in SEQ ID NO. 35 and Figure 4 under the control of the human cytomegalovirus IE enhancer and  
25 promoter and the beta-globin intron and, flanked by 5' and 3' apolipoprotein SAR elements.

In a further embodiment, the expression vector comprises a nucleic acid molecule encoding erythropoietin and having the sequence shown in SEQ ID NO. 35 and Figure  
30 4, flanked by 5' and 3' apolipoprotein SAR elements under the control of elongation factor - 1 alpha promoter and intron.

The present invention still further provides a mammalian cell stably transfected with the expression  
35 vector of the invention. The mammalian cell may be any mammalian cell, for example CHO-K1, BHK, Namalwa. An aspect of the invention provides a mammalian cell, lacking



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multiple copies of an amplified selectable marker gene and capable of expressing recombinant EPO in vitro at levels of at least 1,500, preferably over 2,000, most preferably from 2,000 to 10,000 u/10<sup>6</sup> cells in 24 hours.

5 In yet a further aspect, the invention provides a method of expressing recombinant mammalian erythropoietin comprising the steps of culturing a transfected mammalian cell of the invention in a suitable medium until sufficient amounts of erythropoietin are produced by the  
10 cell and separating the erythropoietin produced.

The present invention also relates to a method of preparing recombinant erythropoietin comprising transfecting a mammalian cell with an expression vector comprising a nucleic acid molecule encoding mammalian  
15 erythropoietin, an expression control sequence operatively linked thereto and at least one SAR element; culturing the transfected cell in a suitable medium until sufficient amounts of erythropoietin are produced by the cell and separating the erythropoietin produced. In a preferred  
20 embodiment, the erythropoietin is produced at levels of at least 2,000, most preferably from 2,000 to 10,000 u/10<sup>6</sup> cells in 24 hours in the absence of gene amplification. The invention also relates to erythropoietin produced by the method of the invention.

25 In an embodiment of the method, the mammalian cell is further transfected with a selectable marker gene and transfected cells are selected in conditions where the activity of the product encoded by the selectable marker gene is necessary for survival of the cells. In a  
30 preferred embodiment, the selectable marker gene is pSV2neo.

In a further embodiment, the method comprises the additional step of identifying and selecting cells producing high levels of erythropoietin; cloning the  
35 selected cells; establishing long term cell lines from the selected cells and; culturing the selected cells in a suitable medium until sufficient amounts of erythropoietin

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are produced by the cell and separating the erythropoietin produced.

The present invention further provides a transgenic non-human animal or embryo whose germ cells and somatic  
5 cells contain a DNA construct comprising the recombinant DNA molecule of the invention.

#### **DESCRIPTION OF THE DRAWINGS**

The invention will be better understood with reference to the drawings in which:

10 Figure 1A is a schematic representation of the procedure for synthesizing the erythropoietin gene;

Figure 1B shows the DNA sequences of the oligonucleotides synthesized to construct the erythropoietin gene;

15 Figure 2A is a schematic representation of the procedure for synthesizing the EPO<sub>short</sub> sequence;

Figure 2B is a schematic representation of the procedure for synthesizing the EPO<sub>long</sub> sequence;

20 Figure 3A shows the DNA sequence of the erythropoietin gene;

Figure 3B shows the amino acid sequence of the mature erythropoietin protein;

Figure 4 shows the DNA sequence of the EPO<sub>long</sub> sequence;

25 Figure 5 shows the DNA sequence of the 3' SAR element of human apolipoprotein B;

Figure 6 shows the DNA sequence of Rh10;

Figure 7 shows a restriction map of the vector pLW18 and;

30 Figure 8 shows a restriction map of the vector pLW19.

#### **DETAILED DESCRIPTION OF THE INVENTION**

As hereinbefore noted, the present inventor has significantly found that recombinant EPO expression is  
35 increased in host cells transfected with recombinant DNA encoding EPO operatively linked to an expression control sequence and SAR elements, compared to host cells

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transfected with EPO in the absence of SAR elements.

The present invention thus provides a recombinant DNA molecule adapted for transfection of a host cell comprising a nucleic acid molecule encoding mammalian erythropoietin, an expression control sequence operatively linked thereto and at least one SAR element. In a preferred embodiment the EPO is human EPO and in a particularly preferred embodiment the nucleic acid molecule has a sequence which encodes erythropoietin having the amino acid sequence as shown in SEQ ID NO 33 and 34 and Figure 3B. In a particular embodiment, the nucleic acid molecule has the sequence as shown in SEQ ID NOS. 35 or 33.

#### **1. Nucleic Acid Molecules Encoding Erythropoietin**

The term "a nucleic acid molecule encoding mammalian EPO" as used herein means any nucleic acid molecule which encodes biologically active EPO. It will be appreciated that, within the context of the present invention, EPO may include various structural forms of the primary protein which retain biological activity. Biologically active EPO will include analogues of EPO having altered activity, for example having greater biological activity than EPO. Biological activity of EPO may be readily determined by the methods referred to herein.

Nucleic acid molecules encoding EPO include any sequence of nucleic acids which encode biologically active EPO, preferably, the nucleic acid molecule also encodes the leader sequence of the prepeptide to permit secretion of EPO from a cell transfected with the recombinant DNA molecule of the invention. The amino acid sequence of the leader sequence of the prepeptide is shown in SEQ ID NO. 33, from amino acid number -27 to -1. In an embodiment, the nucleic acid molecule encodes a peptide having the amino acid sequence as shown in SEQ ID NO. 33 and 34 and Figure 3B. Nucleic acid molecules encoding EPO include the entire EPO gene sequence as shown in SEQ ID NO. 33, one or more fragments of this sequence encoding the EPO

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prepeptide (nucleotides 625-637, 1201-1346, 1605-1691, 2303-2482 and 2617-2772 in SEQ ID NO. 33) or the mature EPO peptide (nucleotides 1269-1346, 1605-1691, 2303-2482 and 2617-2769 in SEQ ID NO. 33) or nucleic acid molecules  
5 having substantial homology thereto, or any fragment thereof encoding biologically active EPO, such as the EPO<sub>long</sub> sequence shown in SEQ ID NO. 35 and Figure 4.

It will be appreciated that the invention includes nucleotide or amino acid sequences which have substantial  
10 sequence homology with the nucleotide and amino acid sequences shown in SEQ ID NOS: 33, 34 and 35 and Figures 3A, 3B and 4. The term "sequences having substantial sequence homology" means those nucleotide or amino acid sequences which have slight or inconsequential sequence  
15 variations from the sequences disclosed in SEQ ID NOS. 33, 34 and 35 i.e. the homologous sequences function in substantially the same manner to produce substantially the same polypeptides as the actual sequences. Due to code degeneracy, for example, there may be considerable  
20 variation in nucleotide sequences encoding the same amino acid sequence. The variations may be attributable to local mutations or structural modifications.

It will also be appreciated that a double stranded nucleotide sequence comprising a DNA segment of the  
25 invention or an oligonucleotide fragment thereof, hydrogen bonded to a complementary nucleotide base sequence, and an RNA made by transcription of this double stranded nucleotide sequence, are contemplated within the scope of the invention.

30 A number of unique restriction sequences for restriction enzymes are incorporated in the DNA sequence identified in SEQ ID NO: 33, and 35 and in Figures 3A, and 4 respectively, and these provide access to nucleotide sequences which code for polypeptides unique to  
35 EPO. DNA sequences unique to EPO or isoforms thereof, can also be constructed by chemical synthesis and enzymatic ligation reactions carried out by procedures known in the

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art.

Mutations may be introduced at particular loci for instance by synthesizing oligonucleotides containing a mutant sequence, flanked by restriction sites enabling  
5 ligation to fragments of the native sequence. Following ligation, the resulting reconstructed sequence encodes a derivative having the desired amino acid insertion, substitution, or deletion.

Alternatively, oligonucleotide-directed site-  
10 specific mutagenesis procedures may be employed to provide an altered gene having particular codons altered according to the substitution, deletion, or insertion required. Deletion or truncation derivatives of EPO may also be constructed by utilizing convenient restriction  
15 endonuclease sites adjacent to the desired deletion. Subsequent to restriction, overhangs may be filled in or removed, and the DNA religated. Exemplary methods of making the alterations set forth above are disclosed by Sambrook et al. (Molecular cloning A Laboratory Manual, 2d  
20 Ed., Cold Spring Harbor Laboratory Press, 1989).

The scope of the present invention also includes conjugates of EPO along with other molecules such as proteins or polypeptides. This may be accomplished, for example, by the synthesis of N-terminal or C-terminal  
25 fusion proteins or fragments of proteins to facilitate purification or identification of EPO (see U.S. Patent No. 4,851,341, see also, Hopp et al., Bio/Technology 6:1204, 1988.) Thus, fusion proteins may be prepared by fusing through recombinant techniques the N-terminal or C-  
30 terminal of EPO or other portions thereof, and the sequence of a selected protein with a desired biological function. The resultant fusion proteins contain EPO or a portion thereof fused to the selected protein or portion thereof. Examples of proteins which may be selected to  
35 prepare fusion proteins include lymphokines such as gamma interferon, tumor necrosis factor, IL-1, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, GM-CSF, CSF-1

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and G-CSF, nerve growth factor, protein A, protein G, GST and the Fc portion of immunoglobulin molecules.

Nucleic acid molecules encoding EPO may be chemically synthesized or may be cloned from a genomic or  
5 cDNA mammalian library using oligonucleotide probes derived from the known EPO sequences following standard procedures. In this manner, nucleic acid molecules encoding EPO may be obtained from the cells of a selected mammal.

10 For example, cDNA sequences encoding mammalian EPO may be isolated by constructing cDNA libraries derived from reverse transcription of mRNA in cells from the selected mammal which express EPO, for example adult kidney cells or foetal liver cells. Increased levels of  
15 expression in the cell may also be achieved, for example by inducing anemia in the mammal. DNA oligonucleotide probes may be used to screen the library for positive clones. Genomic DNA libraries may also be constructed and screened by plaque hybridization using fragments of EPO  
20 cDNA as probes.

Nucleic acid molecules which encode EPO may also be obtained from a variety of sources, including for example, depositories which contain plasmids encoding EPO sequences including the American Type Culture Collection (ATCC,  
25 Rockville Maryland), and the British Biotechnology Limited (Cowley, Oxford England). EPO DNA as described in Lin, (1985, Proc. Natl. Acad. Sci. U.S.A. 82:7580) is deposited as HUMERPA, Accession No M11319.

Various post translational modifications are  
30 contemplated to the EPO encoded by the nucleic acid molecule. For example, EPO may be in the form of acidic or basic salts, or in neutral form. In addition, individual amino acid residues may be modified by, for example, oxidation or reduction. Furthermore, various  
35 substitutions, deletions, or additions may be made to the amino acid or DNA nucleic acid sequences, the net effect of which is to retain biological activity of EPO.

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In a preferred embodiment of the invention, the nucleic acid molecule of the invention comprises the EPO<sub>long</sub> nucleotide molecule encoding EPO, shown in SEQ ID NO. 35 and Figure 4. EPO<sub>long</sub> may be chemically synthesized by assembling short nucleotides, for example, as shown in Figure 1B and SEQ ID NOS. 1 to 32. The procedure for the synthesis and assembly of the EPO<sub>long</sub> sequence from the short nucleotides is shown schematically in Figure 2B.

## 2. SAR Elements

The term "SAR elements" as used herein refers to DNA sequences having an affinity or intrinsic binding ability for the nuclear scaffold or matrix. SAR elements are usually 300 or more base pairs long, require a redundancy of sequence information and contain multiple sites of protein-DNA interaction. Preferably SAR elements are found in non coding regions: in flanking regions or introns.

Suitable SAR elements for use in the invention are those SAR elements which promote elevated and position-independent gene activity in stable transfectants. SAR elements may be obtained, for example, from eukaryotes including mammals, plants, insects and yeast, preferably mammals. SAR elements are preferably selected which co-map with the boundaries of a chromatin domain. SAR elements co-mapping with the chromatin domain boundary are preferred for the recombinant DNA molecules of the invention to promote the formation of an independent domain containing the nucleic acid molecule encoding mammalian EPO to be expressed in stable transfectants.

Examples of preferred SAR elements which co-map with the chromatin domain boundary include the following: the 5' human apoB SAR element (Rh10), a XbaI fragment spanning nucleotides -5,262 to -2,735 of the human apoB gene, as shown in SEQ ID 37 and Figure 6 and as described in Levy-Wilson and Fortier, 1989, (J. Biol. Chem. 264:21196). This region actually contains two SAR elements, a proximal and distal one. It is contemplated that either the

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proximal and distal SAR elements together or the distal element alone may be used. The 3' human apoB SAR element extends between nucleotides +43,186 and +43,850. The DNA sequence of this 665 bp region is described in Levy-Wilson and Fortier, 1989, supra and shown in SEQ ID NO. 36 and Figure 5. A 60-bp deleted 3'human apoB SAR element (Rh32), is also suitable for use in the invention. The deletion is shown within brackets in Figure 5.

Examples of suitable protocols for identifying SAR elements for use in the present invention are described below. The high salt method may be used to identify SAR elements by measuring the ability of labelled naked DNA fragments to bind nuclear matrices in the presence of unlabelled competitor DNA, typically *E. coli* DNA. DNA fragments bound to nuclear matrices under these conditions are operationally defined as SAR elements. The nuclear matrices may be isolated by the 2 M NaCl extraction of DNAase I-digested nuclei (Cockerill and Garrard, 1986, Cell 44, 273-282). Chromosomal loop anchorage of the kappa immunoglobulin gene occurs next to the enhancer in a region containing topoisomerase II sites. (Cockerill, 1990, Nucleic. Acids. Res. 18, 2643-2648).

In the low salt method for identifying SAR elements nuclei are heated to 37°C and then extracted with a buffer containing 25 mM 3,5-diiodosalicylic acid lithium salt (LIS) which removes histones. The LIS-treated nuclei are extracted several times with a low salt buffer. The extracted nuclei are digested with restriction endonucleases. The solubilized (non-matrix bound) DNA fragments are removed by centrifugation. Southern blot hybridization with labelled probes identifies the DNA fragments bound to the nuclear matrix (Mirkovitch et al., 1984, Cell 39, 223-232). Both the above-noted methods yield essentially the same result, that is, a sequence identified as a SAR element with the high salt method will also be identified as a SAR element by the Lis-low salt method.



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The SAR element may be inserted into the recombinant molecule of the invention upstream or downstream from the nucleic acid molecule encoding EPO and the operatively linked expression control sequence. Preferably, the SAR element is inserted within 0.1 to 100 kb upstream or downstream, more preferably from 0.1 to 50 kb, most preferably 0.5 to 10. In a preferred embodiment, more than one SAR element should be inserted into the recombinant molecule of the invention, preferably the SAR elements should be located in flanking positions both upstream and downstream from the nucleic acid molecule encoding EPO and the operatively linked expression control sequence. The use of flanking SAR elements in the nucleic acid molecules may allow the SAR elements to form an independent loop or chromatin domain, which is insulated from the effects of neighbouring chromatin. Accordingly, EPO gene expression may be position-independent and the level of expression should be directly proportional to the number of integrated copies of the recombinant DNA molecules of the invention. Preferably, the SAR elements should be inserted in non-coding regions of the recombinant DNA molecule.

The recombinant DNA molecules of the invention may be advantageously used to express elevated levels of mammalian EPO. Routine procedures may be employed to confirm that the SAR elements selected by the above-noted protocols are useful for expressing elevated levels of mammalian EPO. For example, appropriate expression vectors comprising a nucleic acid molecule encoding mammalian erythropoietin and an expression control sequence operatively linked thereto may be constructed with and without (control vectors) SAR elements. Mammalian host cells may be stably transfected with an EPO expression vector having a selectable marker gene or may be stably co-transfected with an EPO expression vector and a selectable marker gene vector, such as a pSV2-neo vector (which carries the gene conferring the resistance

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to the antibiotic G-418). The levels of EPO secreted may be determined, for example, by RIA and the effect of the flanking SAR elements on EPO production may be determined. Transfected cell populations producing the highest levels of EPO as detected by RIA, may be selected and subjected to successive rounds of cloning by the dilution method.

### **3. Expression Control Sequences**

Suitable expression control sequences may be derived from a variety of mammalian sources. Selection of appropriate regulatory elements is dependent on the host cell chosen, and may be readily accomplished by one of ordinary skill in the art. Examples of regulatory elements include: a transcriptional promoter and enhancer or RNA polymerase binding sequence, splice signals, polyadenylation signals, including a translation initiation signal. Additionally, depending on the host cell chosen and the vector employed, other genetic elements, such as an origin of replication, additional DNA restriction sites, enhancers, sequences conferring inducibility of transcription, and selectable markers, may be incorporated into the expression vector.

Strong promoters (or enhancer/promoters) are preferably selected. A strong promoter is one which will direct the transcription of a gene whose product is abundant in the cell. The relative strength and specificities of a promoter/enhancer may be compared in comparative transient transfection assays. In an embodiment, a promoter may be selected which has little cell-type or species preference and which can therefore be strong when transfected into a variety of cell types.

In preferred embodiments of the invention, the EF1 promoter or the human cytomegalovirus (hCMV) IE (immediate-early) enhancer and promoter may be used. The human EF-1 $\alpha$  gene promoter is stronger than the adenovirus major late promoter in a cell free system (Uetsuki et al., 1989, J. Biol. Chem. 264:5791). The EF1 promoter is the promoter for the human chromosomal gene for the

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polypeptide chain elongation factor-1 $\alpha$ : extending from -303 to -1 for genomic constructs and from -303 to +986 for cDNA constructs. For the cDNA constructs, the EF1 DNA sequence includes the promoter as well as exon 1 and  
5 intron 1 (Uetsuki et al, 1989, J. Biol. Chem. 264, 5791-5798). The human cytomegalovirus (hCMV) IE (immediate-early) enhancer and promoter, extend from -598 to +54 in the sequence (Kay and Humphries, 1991, Methods in Molecular and Cellular Biology 2, 254-265).

10 **4. Methods of Expressing EPO**

As hereinbefore noted, the present invention also provides expression vectors which include the recombinant DNA molecule of the invention and mammalian cells stably transfected with the expression vector.

15 Suitable expression vectors, such as plasmids, bacteriophage, retroviruses and cosmids are known in the art. Many plasmids suitable for transfecting host cells are well known in the art, including among others, pBR322 (see Bolivar et al., Gene 2:9S, 1977), the pUC plasmids  
20 pUC18, pUC19, pUC118, pUC119 (see Messing, Meth in Enzymology 101:20-77, 1983 and Vieira and Messing, Gene 19:259-268, 1982), and pNH8A, pNH16a, pNH18a, and Bluescript M13 (Stratagene, La Jolla, Calif.). Retroviral vectors are reviewed in Eglitis and Anderson, 1988,  
25 Biotechniques 6:608. Suitable expression vectors are those which are stably incorporated into the chromosome of the mammalian host cell.

The recombinant DNA molecule of the invention may be expressed by a wide variety of mammalian cells. Methods  
30 for transfecting such cells to express foreign DNA are well known in the art (see, e.g., Itakura et al., U.S. Patent No. 4,704,362; Hinnen et al., PNAS USA 75:1929-1933, 1978; Murray et al., U.S. Patent No. 4,801,542; Upshall et al., U.S. Patent No. 4,935,349; Hagen et al.,  
35 U.S. Patent No. 4,784,950; Axel et al., U.S. Patent No. 4,399,216; Goeddel et al., U.S. Patent No. 4,766,075; and Sambrook et al. Molecular Cloning A Laboratory Manual, 2nd

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edition, Cold Spring Harbor Laboratory Press, 1989, all of which are incorporated herein by reference). Suitable expression vectors include vectors having a selectable marker gene.

5 Mammalian cells suitable for carrying out the present invention include, among others: COS (e.g., ATCC No. CRL 1650 or 1651), BHK (e.g., ATCC No. CRL 6281), CHO-K1 (ATCC No. CCL 61), HeLa (e.g., ATCC No. CCL 2), 293 (ATCC No. 1573) and NS-1 cells. As noted above, suitable  
10 expression vectors for directing expression in mammalian cells generally include a promoter, as well as other transcriptional and translational control sequences. Common promoters include SV40, MMTV, metallothionein-1, adenovirus Ela, CMV, immediate early, immunoglobulin heavy  
15 chain promoter and enhancer, human cytomegalovirus 1E enhancer and promoter and RSV-LTR. Protocols for the transfection of mammalian cells are well known to those of ordinary skill in the art. Representative methods include calcium phosphate mediated gene transfer, electroporation,  
20 retroviral, and protoplast fusion-mediated transfection (see Sambrook et al., supra).

Given the teachings provided herein, promoters, terminators, and methods for introducing expression vectors of an appropriate type into mammalian cells may be  
25 readily accomplished. Accordingly, the invention also relates to mammalian cells stably transfected with an expression vector of the invention. The term "stably transfected" refers to the fact that suitable expression vectors are those which stably incorporate the recombinant  
30 DNA molecule of the invention into the chromosomes of the mammalian cell.

The invention also relates to a mammalian cell lacking a selectable marker gene capable of being amplified and, capable of expressing recombinant EPO in  
35 vitro at levels of at least 1,500, preferably at least 2,000, most preferably 2,000 to 10,000 u/10<sup>6</sup> cells in 24 hours.

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In an embodiment, the invention also relates to a mammalian cell, having less than 1,000, preferably less than 100, most preferably less than 10 copies of a selectable marker gene and capable of expressing  
5 recombinant EPO in vitro at levels of at least 1,500, preferably at least 2,000, most preferably 2,000 to 10,000 u/10<sup>6</sup> cells in 24 hours.

Amplification refers, for example, to the process of culturing cells in appropriate medium conditions to  
10 select cells resistant to the drug methotrexate. Such cells have been found to be resistant to methotrexate due to an amplification of the number of their gene encoding dihydrofolate reductase (DHFR). Selection for cells resistant to methotrexate produces cells containing  
15 greater numbers of DHFR genes. Passenger genes, such as the EPO gene carried on the expression vector along with the DHFR gene or co-transfected with the DHFR gene may also be increased in their gene copy number. Cells which have been amplified thus have multiple copies of the  
20 selectable marker gene in addition to the passenger gene.

It is an advantage of the present invention that high levels of EPO expression may be achieved without the need for amplification. Thus the mammalian cells of the invention express high levels of EPO and do not express  
25 high levels of a selectable marker gene. The expression of high levels of selectable marker genes in amplified cells may interfere with the cell's ability to remain stable in long term culture and to express high levels of EPO in long term culture. Amplified cells may carry  
30 certain disadvantages such as non-specific toxicity associated with exposure of the cells to the inhibitory drug or compound.

Selectable marker genes which may be amplified in a mammalian host cell are known in the art and include the  
35 genes encoding proteins conferring resistance to chloramphenicol (Wood et al., CSHSQB 51:1027, 1986), methotrexate (Miller, MC Biol., 5:431, 1985; Corey et al.,

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Blood 75:337, 1990; Williams et al., Proc. Natl. Acad. Sci. USA, 83:2566, 1986; Stead et al., Blood 71:742, 1988), mycophenolic acid (Stuhlmann et al., Proc. Natl. Acad. Sci. USA 81:7151, 1984), or various chemotherapeutic  
5 agents (Guild et al., Proc. Natl. Acad. Sci USA 85:1595, 1988; Kane et al., Gene 84:439, 1989; Choi et al., Proc. Natl. Acad. Sci. USA; Sorrentino, et al., Science 257:99, 1992). A discussion of selectable marker genes, including those capable of being amplified is provided in Kreigler,  
10 1990, "Gene Transfer and Expression, A Laboratory Manual", Chapter 6, Stockton Press, and include the gene encoding dihydrofolate reductase.

Within another aspect, the present invention relates to a method of preparing recombinant erythropoietin  
15 comprising transfecting a mammalian cell with an expression vector comprising a nucleic acid molecule encoding mammalian erythropoietin, an expression control sequence operatively linked thereto and at least one SAR element; culturing the transfected cell in a suitable  
20 medium until sufficient amounts of erythropoietin are produced by the cell and separating the erythropoietin produced. In an embodiment, the mammalian cell may be further transfected with a selectable marker gene and the transfected cells may be selected by means of the  
25 selectable marker gene. Examples of selectable marker genes are given above and include neo.

In an embodiment of the method, cells producing high levels of erythropoietin may be identified and selected and subcloned to establish long term cell lines from the  
30 selected cells and; the selected cells may be cultured in a suitable medium until sufficient amounts of erythropoietin are produced by the cell. EPO produced may then be separated from the medium.

It is an advantage of the method of the invention  
35 that cell lines may be established which are stable over the long term, at least over six months. The long term cell lines of the invention express consistently high

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levels of EPO and may be maintained without the selective pressure often required to maintain the high copy number of amplified genes in cell lines which have been subjected to amplification.

5 EPO may be prepared by culturing the host/vector systems described above, in order to express the EPO. Recombinantly produced EPO may be further separated and further purified as described in more detail below.

Biologically active EPO expressed may be assayed by  
10 known procedures such as tritiated thymidine uptake by mouse spleen cell erythrocyte precursors (Krystal, et al., 1986, Blood 67:71); the exhypoxic mouse method using <sup>59</sup>Fe incorporation into erythrocyte precursors (Cotes and Bangham, 1961, Nature 191:1065); <sup>59</sup>Fe uptake into fetal  
15 mouse liver cells (Dunn et al, 1975, Exp. Hematol. 3:65); and the starved rat method (Goldwasser and Gross, 1975, Methods Enzymol. 37:109). EPO once expressed may also be quantitated, for example by RIA, separated and purified by known techniques such as ultrafiltration, flat-bed  
20 electrofocusing, gel filtration, electrophoresis, isotachophoresis and various forms of chromatography, such as ion exchange, adsorption chromatography, column electrophoresis and various forms of HPLC. Procedures for the chromatographic separation of EPO are described, for  
25 example in U.S. Patent No 4,667,016.

The present invention also relates to transgenic non-human mammals or embryos whose germ cells and somatic cells contain a DNA construct comprising the recombinant DNA molecule of the invention. The recombinant DNA  
30 molecule of the invention may be expressed in non-human transgenic animals such as mice, rats, rabbits, sheep, cows and pigs (see Hammer et al. (Nature 315:680-683, 1985), Palmiter et al. (Science 222:809-814, 1983), Brinster et al. (Proc Natl. Acad. Sci USA 82:44384442, 1985), Palmiter and Brinster (Cell. 41:343-345, 1985) and  
35 U.S. Patent No. 4,736,866). Briefly, an expression unit, including a DNA sequence to be expressed together with

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appropriately positioned expression control sequences, is introduced into pronuclei of fertilized eggs. Introduction of DNA is commonly done by microinjection. Integration of the injected DNA is detected by blot  
5 analysis of DNA from tissue samples, typically samples of tail tissue. It is preferred that the introduced DNA be incorporated into the germ line of the animal so that it is passed on to the animal's progeny. Tissue-specific expression may be achieved through the use of a tissue-  
10 specific promoter, or through the use of an inducible promoter, such as the metallothionein gene promoter (Palmiter et al., 1983, *ibid*), which allows regulated expression of the transgene. Alternatively, yeast  
15 artificial chromosomes (YACs) may be utilized to introduce DNA into embryo-derived stem cells by fusion with yeast spheroblasts carrying the YAC (see Capecchi, *Nature* 362:255-258, 1993; Jakobovits et al., *Nature* 362:255-258, 1993). Utilizing such methods, animals may be developed which express EPO in tissues. Tissue specific promoters  
20 may be used to target expression of EPO in cells. Tissue specific promoters include the 5' or 3' flanking sequences of the beta-globin, elastase, alpha-fetoprotein, alpha-A crystalline, an erythroid specific transcriptional element and insulin genes (Yee, et al. (1989) *P.N.A.S., U.S.A.*  
25 86, 5873-5877; Swift, et al. 1984, *Cell* 38:639; Storb et al., *Nature (Lond.)* 310:238; Grosschedl et al., 1985 *Cell* 41:885; Shani, 1985 *Nature (Lond)* 314:238 and Chada et al, 1985, *Nature (Lond)*). The use of SAR elements in the development of transgenic animals is described for example  
30 in Xu, 1989, *J. Biol. Chem.* 264:21190; McKnight et al., 1992, *Proc. Natl. Acad. Sci. U.S.A.* 89:6943; Brooks, et al., 1994, *Mol. Cell. Biol.* 14:2243 and; Forrester, et al., 1994, *Science* 265:1221.

In a preferred embodiment suitable promoters and/or  
35 enhancers may be selected from mammary gland specific genes which are normally only expressed in milk, for example the genes encoding  $\alpha$ -casein (Gene Pharming,



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Leiden, Netherlands),  $\beta$ -casein (Genzyme Transgenics Corp. Framingham, Mass.),  $\gamma$ -casein,  $\kappa$ -casein  $\alpha$ -lactalbumin  $\beta$ -lactalbumin  $\beta$ -lactoglobulin (PPL Therapeutics Ltd, Edinburgh, Scotland) and whey acidic protein (Altra Bio Inc., Arden Hills MN). Methods for targeting recombinant gene expression to the mammary gland of a mammal are described, for example, in U.S. Patent No. 5,304,489. Briefly, a DNA construct with the recombinant DNA molecule of the invention comprising a mammary gland specific promoter may be microinjected into a newly fertilized egg leading to integration of the construct into the genome and secretion of the protein into the milk of a mature lactating female. EPO expressed in the milk may be purified from the milk using standard procedures after skimming off the milk fat, separating out the caseins and precipitating EPO out of the whey fraction, followed by standard protein purification procedures for EPO as described elsewhere herein.

A major problem in the generation of transgenic mammals for the expression of recombinant proteins has been the varying levels of expression which result due to chromosomal factors in the local environment where the construct integrates, for example regulatory elements and the state of the chromatin (open or closed). As a result, generating a transgenic mammal that produces high levels of a recombinant protein has been achieved only by laborious trial and error.

The present invention provides a transgenic non-human animal whose germ cells and somatic cells contain a DNA construct comprising the recombinant DNA molecules of the invention, comprising a tissue specific promoter, preferably the tissue specific promoter is a promoter which specifically expresses EPO in milk. In a particularly preferred embodiment, the recombinant molecule contained in the transgenic non-human mammal comprises the nucleic acid molecule encoding EPO and the expression control sequence operatively linked thereto,

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flanked by SAR elements. An advantage of DNA constructs having flanking SAR elements is that expression of EPO may be independent of the site of integration of the construct as the construct is insulated from surrounding chromosome material by the SAR elements, which define an open chromatin domain.

#### 5. Applications

It will be apparent that the recombinant DNA molecules, expression vectors, transfected host cells, methods and transgenic animals of the invention will be useful for the efficient expression and production of recombinant mammalian EPO in vitro and in vivo as described above.

Recombinant EPO may be used to treat animals and human patients, including patients having anemia as a result of chronic renal failure. As EPO is the primary regulator of red blood cell formation, it has applications in both the diagnosis and treatment of disorders of red blood cell production and has potential applications for treating a range of conditions such as anemia, sickle cell disease, conditions where red cells are depleted (for example in bone marrow transplants), thalassemia, cystic fibrosis, menstrual disorders, acute blood loss and conditions involving abnormal erythropoiesis (for example cancers of the haemopoietic system), conditions involving destruction of red blood cells by over exposure to radiation, reduction in oxygen intake at high altitudes, complications or disorders secondary to AIDS and prolonged unconsciousness.

The recombinant DNA molecules, expression vectors and transformed mammalian cells of the invention will also have useful applications in gene therapy, whereby a functional EPO gene is introduced into a mammal in need thereof, for example mammals having anemias. The transfer of the recombinant DNA molecule of the invention into mammalian cells may be used, for example in gene therapy to correct an inherited or acquired disorder through the

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synthesis of missing or defective EPO gene products in vivo.

The recombinant DNA molecule of the invention may be used in gene therapy as briefly described below. The recombinant DNA molecule may be introduced into cells of a mammal, for example haemopoietic stem cells removed from the bone marrow or blood of the mammal. Hemopoietic stem cells are particularly suited to somatic gene therapy as regenerative bone marrow cells may be readily isolated, modified by gene transfer and transplanted into an immunocompromised host to reconstitute the host's hemopoietic system. Suitable hemopoietic stem cells include primitive hemopoietic stem cells capable of initiating long term culture (Sutherland et al., Blood, Vol. 74, p. 1563, 1986 and Udomsakdi et al., Exp. Hematol., Vol. 19, p. 338, 1991.) Suitable cells also include fibroblasts and hepatocytes.

The recombinant DNA molecules of the invention may be introduced into the cells by known methods, including calcium phosphate mediated transfection described herein or retroviral mediated uptake. The recombinant DNA molecule of the invention may be directly introduced into cells or tissues in vivo using delivery vehicles such as retroviral vectors, adenoviral vectors and DNA virus vectors. They may also be introduced into cells in vivo using physical techniques such as microinjection and electroporation or chemical methods such as coprecipitation and incorporation of DNA into liposomes. Recombinant molecules may also be delivered in the form of an aerosol or by lavage. The recombinant DNA molecules of the invention may also be applied extracellularly such as by direct injection into cells. Freed et al., New Eng. J. Med. 327(22):1549-1555, 1992, describe a method for injecting fetal cells into brains of Parkinson's patients. Gene therapy involving bone marrow transplant with recombinant primary hemopoietic stem cells requires efficient gene transfer into the stem cells. As a very

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small number of primary stem cells can reconstitute the entire host hemopoietic system it is important that the transferred gene be efficiently expressed in the recombinant stem cells transferred. Thus it is expected  
5 that the recombinant molecules of the invention will be particularly advantageous for use in gene therapy to correct defects in the erythropoietin gene.

As hereinbefore noted, the recombinant DNA molecules of the invention, having flanking SAR elements, are  
10 particularly useful for the expression of EPO in transgenic mammals and therefore they are particularly useful in gene therapy as the SAR elements define an open chromatin domain and insulate the construct from the surrounding chromosome material, thereby, providing  
15 position-independent expression.

The following non-limiting examples are illustrative of the present invention:

#### EXAMPLES

##### EXAMPLE 1

##### 20 Synthesis of EPO cDNA Sequences

##### EPO Oligonucleotide synthesis

EPO oligonucleotides were synthesized with the Applied Biosystems Inc. 392 DNA/RNA Synthesizer at the 0.2  $\mu$ m scale (~500  $\mu$ g). Oligonucleotide 5' ends were  
25 phosphorylated, except for the ones coinciding with the 5' ends of each of the four blocks. Oligonucleotides lacking 5' phosphorylation were purified using the Applied Biosystems Oligo Purification Cartridge (OPC). Oligonucleotides with 5' phosphorylation were purified  
30 from acrylamide gels (according to the protocol described in Sambrook, Fritsch, and Maniatis, 1989. Molecular Cloning: A Laboratory Manual, 2nd Edition. Cold Spring Harbor Laboratory Press, pp. 11.23 - 11.30).  $C_{18}$  Sep-Pak cartridges from Millipore were used to remove salts. All  
35 oligos were resuspended in double distilled  $H_2O$ .

Oligonucleotides EPO1 (SEQ ID NO. 1), EPO2 (SEQ ID NO. 2), EPO3 (SEQ ID NO. 3), EPO4 (SEQ ID NO. 4), EPO5

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(SEQ ID NO. 5), EPO6 (SEQ ID NO. 6), EPO7 (SEQ ID NO. 7), EPO8 (SEQ ID NO. 8), EPO9 (SEQ ID NO. 9), EPO10 (SEQ ID NO. 10), EPO11 (SEQ ID NO. 11), EPO12 (SEQ ID NO. 12), EPO13 (SEQ ID NO. 13), EPO13b (SEQ ID NO. 14), EPO14 (SEQ ID NO. 15), EPO15 (SEQ ID NO. 16), EPO1 $\alpha$  (SEQ ID NO. 17), EPO2 $\alpha$  (SEQ ID NO. 18), EPO3 $\alpha$  (SEQ ID NO. 19), EPO4 $\alpha$  (SEQ ID NO. 20), EPO5 $\alpha$  (SEQ ID NO. 21), EPO6 $\alpha$  (SEQ ID NO. 22), EPO7 $\alpha$  (SEQ ID NO. 23), EPO8 $\alpha$  (SEQ ID NO. 24), EPO9 $\alpha$  (SEQ ID NO. 25), EPO10 $\alpha$  (SEQ ID NO. 26), EPO11 $\alpha$  (SEQ ID NO. 27), EPO12 $\alpha$  (SEQ ID NO. 28), EPO13 $\alpha$  (SEQ ID NO. 29), EPO13B $\alpha$  (SEQ ID NO. 30), EPO14 $\alpha$  (SEQ ID NO. 31) and EPO15 $\alpha$  (SEQ ID NO. 32) as shown in Figure 1B were synthesized. All oligonucleotides labelled with " $\alpha$ " are for the complementary strand (i.e. negative sense of the EPO gene). Oligonucleotides EPO1 (SEQ ID NO. 1), EPO1 $\alpha$  (SEQ ID NO. 17), EPO15 (SEQ ID NO. 16) and EPO15 $\alpha$  (SEQ ID NO. 32) contain extra bases at their 5'-ends and 3'-ends, respectively, which facilitate construction of a HindIII recognition site.

#### 20 Assembly of blocks 1, 2, 3 and 4

Blocks 1, 2, 3, and 4 as shown in Figure 1A were synthesized by ligating the above-noted EPO oligonucleotides according to the following protocol.

40 pmoles of each oligonucleotide were mixed in a microcentrifuge tube in a final volume of 50 $\mu$ l. The oligonucleotides were annealed by heating at 98°C for 5 minutes in a heat block. The heat block containing the tube was removed from the heating unit and allowed to cool on the benchtop to 30°C (approximately 1.5 hrs). 6 $\mu$ l of 10X T<sub>4</sub> DNA ligase buffer (0.5M Tris-HCl, pH 7.8, 0.1M MgCl<sub>2</sub>, 0.1M DTT, 10mM ATP and 250 $\mu$ g/ml BSA), was added to 2 $\mu$ l (or 2 units) of T<sub>4</sub> DNA ligase, and brought to volume to 60 $\mu$ l with double distilled H<sub>2</sub>O and incubated overnight at 14°C. The mixture was heated at 75°C for 10 minutes to inactivate ligase and then cooled on ice to dissociate non ligated oligonucleotides and ethanol precipitate.

Blocks 1, 2, 3 and 4 were purified as follows.

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Each ligation mix was run on a 3% low melting agarose gel. No band of the expected size was visible on the gel. However, DNA was extracted from the region of the gel where the gene block was expected to be. The rationale was that there was a small amount of the correct gene block present.

The DNA extracted from the agarose gel was amplified by the polymerase chain reaction (with the NE Biolabs Vent DNA polymerase, 1 cycle 30 sec at 98°C, 25-30 cycles 30 sec at 98°C, 40 sec at 50-55°C and 2 min at 72°C, 1 cycle 10 min at 72°C, and cooled at 6°C. The primers were complementary to the ends of the gene blocks and contained a few extra bases so that the entire recognition sites for the specific restriction enzymes flanking each block would be present in each complete block. Blocks were then cloned into the SmaI site of pUC18 or pUC19 plasmids.

Assembled gene blocks were sequenced by the dideoxy-terminator method (Sanger, and Coulson, 1975, J. Mol. Biol. 94, 441-448). Typically, 5-12 clones had to be sequenced for each gene block in order to identify one with the expected sequence.

EPO cDNA sequences were assembled as described in Example 2 herein. EPO<sub>short</sub> and EPO<sub>long</sub> cDNA were assembled into pUC18 as shown in Figures 2A and 2B, to generate pLW20 and pLW21, respectively.

## EXAMPLE 2

### Assembly of Expression Vectors

#### Synthesis and assembly of EPO cDNA

Two EPO cDNA sequences were chemically synthesized: EPO<sub>short</sub> and EPO<sub>long</sub>. The reason for making both a long and a short version of the gene was based on previous results while expressing another gene in COS cells. It had been noted that reducing the length of the 3' non-coding sequence coincided with increased expression of the gene. However, this was not observed with EPO. Therefore the constructs used for this example contained the EPO<sub>long</sub> cDNA,

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as shown in SEQ ID NO. 35 and Figure 4.

Figure 1A, provides a schematic representation of the synthesis and assembly of the EPO<sub>long</sub> cDNA sequence (SEQ ID NO. 35 and Figure 4) from the oligonucleotides. Briefly, the coding and complementary strands from contiguous oligonucleotides which had an average length of 60 bases were synthesized. The breakpoints between the oligonucleotides were chosen such that when two complementary oligonucleotides annealed, cohesive ends compatible with those of the adjoining oligonucleotide pair were created. The EPO<sub>long</sub> cDNA (SEQ ID NO. 35 and Figure 4) was constructed from fifteen oligonucleotide pairs, shown in Figure 1B and SEQ ID NOS. 1-32) The coding strand oligonucleotides were numbered from 1 to 15 except for 13 which was replaced with 13b, and the complementary strand oligonucleotides were numbered 1α to 15α except for 13α which was replaced with 13bα.

The fifteen oligonucleotide pairs were assembled in four blocks. Each junction between adjacent blocks constitutes a unique restriction site. Only three silent mutations were introduced into the EPO<sub>long</sub> cDNA sequence for the creation of unique restriction sites: (1) C at position 22 was replaced with A, to add a BstXI site, (2) C at position 256 was replaced with T, to remove a PstI site, and (3) C at position 705 was replaced with T, to add a SspI site. The assembly of the four blocks was done in PUC18, leading to the vector pLW21. pLW21 is the pUC18 vector with the 788 bp EPO<sub>long</sub> cDNA sequence, as shown in SEQ ID NO. 35 and Figure 4, inserted in the HindIII restriction site.

#### Isolation of genomic EPO DNA

A human leukocyte genomic library was purchased from Clontech and was screened by plaque hybridization with fragments of EPO cDNA as probes. A 2.4kb (2,365bp) EPO genomic clone was isolated. This clone spans the HUMERPA sequence (Genbank, accession number M11319) (SEQ ID NO. 33 and Figure 3A) from nucleotide 499 to nucleotide 2365,

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that is to say its 5' end maps 126 nucleotides upstream of the ATG initiation coding of the EPO pre-protein. The rationale for isolating the EPO gene was that the length of the DNA between the SAR elements would be longer and might allow for an increased effect of the SAR elements.

#### SAR elements

Two human apolipoprotein B (apoB) SAR elements were used which co-map with the boundaries of the human apolipoprotein B gene chromatin domain (Levy-Wilson & Fortier, 1989, J. Biol. Chem. 264:21196). The following clones were used: Rh10 carrying the distal 1212 bp-long 5' human apoB SAR element and 1317 bp of proximal sequence (SEQ ID NO. 37 and Figure 6) and a clone (Rh32) carrying the 605 bp long 3' hu apoB SAR element (SEQ ID NO. 36 and Figure 5).

The DNA sequence of Rh10 was determined by dideoxy-terminator method (Sanger, F. and Coulson, A.R., 1975, J. Mol. Biol. 94:441) and is shown in SEQ ID NO. 37 and Figure 6. The 2529 bp Rh10 sequence consists of the 1212 bp 5' distal human apoB SAR elements and the 1317 bp 5' proximal sequence in the 5' to 3' orientation. The DNA sequence of Rh32 was also determined, and was found to be identical to the sequence published in Levy-Wilson & Fortier (1989, J. Biol. Chem. 264:21196) (SEQ ID NO. 36 and Figure 5), except for a 60 base pair deletion spanning nucleotides 259 to 318, shown within brackets in Figure 5. Contrary to Rh10 which is not a typical SAR sequence, Rh32 is A/T rich and contains 22 copies of the ATATTT motif.

#### Regulatory Elements of Expression Vectors

Two basic expression vectors were derived from pAX111 and pAX142 (renamed pLW19 or pSB3 and pLW18 or pSB2, respectively). pAX111 (Kay & Humphries, Methods Mol. Cell. Biol. 2:254, 1991) carries the human cytomegalovirus IE enhancer and promoter and the  $\beta$ -globin intron, while pAX142 carries the elongation factor-1 $\alpha$  promoter and intron. For each of these vectors there are single restriction sites located on each site of the EPO



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transcription unit, where the SAR elements were introduced (SpeI or EcoRV at the 5' end and HpaI at the 3' end). Maps of the two vectors pLW18 (pSB2) and pLW19 (pSB3) are shown in Figures 7 and 8 respectively.

## 5 Assembly of Expression Vectors

The EPO<sub>long</sub> cDNA sequence (SEQ ID NO.35, Figure 4) or the genomic EPO DNA (SEQ ID NO. 33, Figure 3A) were introduced clockwise into the cloning sites of the expression vectors. In the case of the genomic clones, the  $\beta$ -globin and EF1 introns were removed from the pSB3 and pSB2 vectors, respectively. A summary of the expression vectors is shown in Table 1.

### Construction of pLW24 and pLW25 Vectors

The EPO<sub>long</sub> cDNA (SEQ ID NO.35, Figure 4) was removed from pLW21 by digestion with HindIII, was blunt-ended with the T4 DNA polymerase and was introduced in the clockwise orientation into the SmaI site of pLW18 and pLW19 to generate pLW24 and pLW25, respectively.

### Construction of p24MAR1 and p25MAR1

Introduction of the 5' human apoB SAR element at the 5' end of the EPO transcription unit was accomplished as follows. The 2,529 bp XbaI fragment from Rh10 (SEQ ID NO. 37, Figure 6) was introduced in the clockwise orientation into the SpeI site of pLW24 and pLW25 to generate p5MAR24 and p5MAR25, respectively (Note that XbaI and SpeI sites are compatible).

Introduction of the 60 bp-deleted 3' human apoB SAR element at the 3' end of the EPO transcription unit was accomplished as follows. The 605 bp DraI-RsaI fragment from the plasmid 12 DI Eco was added XhoI linkers and cloned into the XhoI site of the Stratagene pSBII SK(+) plasmid, to generate plasmid 3'apoBX (Rh32) (SEQ ID NO. 36, Figure 5). The ~605 bp SAR fragment was then removed from Rh32 by XhoI digestion, blunt-ended with the Klenow enzyme and cloned into the HpaI site of p5MAR24 and p5MAR25 to generate p24MAR-1 and p25MAR-1, respectively. The orientation of the inserted fragments has not yet been

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determined.

Construction of pAP142 and pAP140

Plasmids pSB2 and pSB3 were derived from pLW18 and pLW19, respectively. A 12bp DNA linker with the EcoRV and SphI restriction sites was inserted into the KpnI site of pLW18 and pLW19 to generate pSB2 and pSB3, respectively. This allowed for introduction of SAR elements into the blunt-end EcoRV restriction site, just upstream of the EPO transcription unit.

10       The EPO<sub>long</sub> cDNA sequence was removed from pLW24 by digestion with Sall and cloned in the clockwise orientation into the Sall site of pSB2 to generate pAP13. pAP5 was generated by removing the EPO<sub>long</sub> cDNA sequence from pLW25 by digestion with BamHI and cloning in the clockwise orientation into the BamHI site of pSB3 to generate pAP5.

20       The Rh32 ~605 bp XhoI SAR fragment (60 bp-deleted 3' human apoB SAR element) was blunt-ended with the T4 DNA polymerase and inserted in the clockwise orientation into the EcoRV site of pAP13 and pAP5 to generate pAP138 and pAP136, respectively. The same Rh32 ~605 bp XhoI blunt-ended SAR fragment was inserted in the clockwise orientation into the HpaI site of pAP138 and pAP136 to generate pAP142 and pAP140, respectively.

25       Construction of pAP59 and pAP67

pAP42 is an intronless version of pSB2. pSB2 was amplified by the polymerase chain reaction between nucleotides 1308 and 321, and the amplified fragment was ligated on itself to produce pAP42. pAP43 is an intronless version of pSB3. The SmaI - XmnI fragment (nucleotides 677 - 1320) was removed from pSB3 to produce pAP43.

35       The EPO genomic clone was isolated from a Clontech library (Cat. No. HL1006d; Lot No. 19412) by probing with the BamHI EPO<sub>short</sub> cDNA fragment from pAP4. The EPO<sub>short</sub> cDNA fragment used as a probe to isolate the genomic EPO clone was a BamHI fragment from pAP4. As for EPO<sub>long</sub> cDNA, EPO<sub>short</sub> cDNA was first assembled into the HindIII site of the

- 33 -

pUC18 vector, to generate the pLW20 vector, blunt-ended with the T4 DNA polymerase and inserted in the clockwise orientation into the SmaI site of pLW19 to generate pLW23. Then the EPO<sub>short</sub> cDNA sequence was removed from pLW23 by  
5 BamHI digestion and cloned in the clockwise orientation into the BamHI site of pSB3 to generate pAP4.

Plaque hybridization was performed according to the method described in "Molecular Cloning. A Laboratory Manual, 2nd Edition. 1989". Edited by Sambrook, Fritsch,  
10 and Maniatis, Cold Spring Harbor Laboratory Press. pp 2.108 - 2.121. A 7.5 kb BamHI fragment (cloned in the Stratagene pBS (KS+) plasmid) was identified as an EPO genomic clone. The identity of this clone was confirmed by polymerase chain reaction between primers EPOGEN 1  
15 (nucleotides 499- 518 in HUMERPA sequence) and EPOGEN 12 (nucleotides 2848 - 2863, on complementary strand, in HUMERPA sequence). This resulted in the expected fragment. The 2,365 bp amplified fragment was cloned in the clockwise orientation into the HincII site of pUC19 to  
20 generate pAP41. The EPO genomic sequence was then removed from pAP41 by EcoRI and HindIII digestion, blunt-ended with the T4 DNA polymerase and inserted in the clockwise orientation into the SmaI site of pAP42 to generate pAP59 or into the blunt-ended BamHI site of pAP43 to generate  
25 pAP67.

#### Construction of pAP123, pAP127, pAP132 and pAP134

The Rh32 ~605 bp XhoI SAR fragment (60 bp-deleted 3' human apoB SAR element) was blunt-ended with the T4 DNA polymerase and inserted in the clockwise orientation into  
30 the HpaI site of pAP59 and pAP67 to generate pAP117 and pAP119, respectively. The same Rh32 ~605 bp XhoI blunt-ended SAR fragment was inserted in the clockwise orientation into the EcoRV site of pAP117 and pAP119 to generate pAP123 and pAP127, respectively.

35 The 2,529 bp XbaI fragment from Rh10 was blunt-ended with the T4 DNA polymerase, and introduced in the clockwise orientation into the EcoRV site of pAP117 and

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pAP119 to generate pAP132 and pAP134, respectively.

### EXAMPLE 3

#### Transfection and Expression of EPO

##### Transfection

5 Briefly as described in more detail below CHO-K1 cells were co-transfected by the calcium phosphate precipitate method by two pairs of vectors: pSV2-neo carrying the resistance to the selective agent G-418 and one of the two following vectors expressing the EPO cDNA  
10 from the EF1 promoter: pLW24 or p24-MAR. pLW24 had no SAR element, while p24-MAR had the 5'apo B SAR element (Rh10) upstream of the EF 1 promoter region and the 60 bp-deleted 3'apoB SAR element (Rh32) downstream of the EPO cDNA.

G-418 was added to the medium to select cells  
15 transfected with pSV2-neo. At least 80% of these transfectants were expected to have been co-transfected with the EPO vector.

The following protocol was used for transfection with the Mammalian Transfection Kit by Stratagene (Catalog  
20 #200285). On day -1 100mm culture dishes were inoculated with exponentially growing mammalian cells at a concentration of  $5 \times 10^4$  cells per ml in 10 ml. The cells were grown overnight at 37°C with the appropriate level of CO<sub>2</sub> and in appropriate medium. For CHO-K1 cells Ham's F12  
25 complete medium with 10% fetal calf serum (FCS) was used and incubation was carried out with 5% CO<sub>2</sub>. The cells were approximately 10-20% confluent on day zero.

The optimal amount of DNA to be used for transfection varied depending on the cell type being used  
30 for transfection. Usually, 10-30 µg of plasmid DNA was used. The plasmid used for selection was generally added at some ratio to the expression vector. For pSV2neo, a ratio of 1:10 to 1:15 was found to be appropriate.

The desired amount of DNA was diluted to 450 µl with  
35 double distilled H<sub>2</sub>O. 50 µl of solution 1 (2.5M Ca Cl<sub>2</sub>, included in the Stratagene kit) was added slowly and

- 35 -

dropwise. 500  $\mu$ l of Solution 2 (2 x BBS, pH 6.95 [consists of 50 mM n, n-bis (2-hydroxyethyl)-2-aminoethanesulfonic acid and buffered saline, 280 mM NaCl and 1.5 mM  $\text{Na}_2\text{HPO}_4$ ]) was added and mixed gently and slowly. It was found to be important to perform these two additions slowly and gently. The mixture was allowed to incubate at room temperature for 10-20 minutes. The precipitate was gently mixed to ensure adequate suspension. The precipitate was added to the culture dropwise while gently swirling the plate to distribute the suspension evenly, followed by incubation for 12-24 hours.

Where spare incubator space was available, it was determined that transfection efficiency could be improved 2-3 times by using lower  $\text{CO}_2$  concentrations at this point (2-4% is recommended; 3% seemed to work well with CHO cells). Normal  $\text{CO}_2$  concentrations were resumed after removal of precipitate on Day 1.

On day 1 the medium was removed by aspiration and the culture rinsed twice with sterile PBS (Phosphate buffered saline without  $\text{Ca}^{2+}$ ,  $\text{Mg}^{2+}$ ) or medium without serum. Fresh complete medium was applied (as on Day - 1) and the cells incubated under optimal  $\text{CO}_2$  concentration for 24 hours.

The cells were split at a ratio ranging from 1:30 to 1:100 into 96 well plates (0.2 ml per well). For CHO-K1 cells, Ham's F12 complete medium with 20% FCS was used at this step. Incubation was carried out at 37°C with 5%  $\text{CO}_2$ .

On day 4 the medium was aspirated and selective medium was added. For transfections using CHO-K1 cells and pSV2neo as the selective plasmid, Ham's F12 complete medium with 20% FCS + 270-400  $\mu$ g/ml G418 was used.

On day 5 or 6 (preferably day 5) and on day 8, the cultures were refed with selective medium.

On day 10, spent medium was collected for RIA assays and frozen. The spent medium was replaced with selective medium.

From day 11 RIA assays, were performed to determine

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levels of EPO production in the medium, using undiluted samples. Samples producing high levels of EPO were cloned and subcultured in Ham's F12 complete medium with 10% FCS when they reached confluency.

#### 5 EPO Production

Levels of EPO production in the medium were determined by RIA as follows.

The following materials and equipment were used in the assay: phosphate RIA buffer: 0.05M  $\text{NaPO}_4$ ; 0.2% BSA; 10 0.02%  $\text{NaN}_3$ , pH 7.4; Erythropoietin (standards, Boehringer Mannheim, Cat.# 1120166, 250 U/ml);  $^{125}\text{I}$ -EPO (Amersham Cat.# IM.219); anti-rabbit IgG (whole molecule) developed in goat, whole antiserum (Sigma Cat.# R-5001 (abbrev. Ab2) polyethylene glycol 8000 (BDH Cat.#B80016); normal rabbit 15 serum (abbrev. NRS); Rabbit anti-EPO, polyclonal, 1mg (abbrev. Ab1) (R&D Systems Cat.# AB-286-NA) and; gamma counter (LKB-Wallac RIAGamma 1272).

The stock concentration of EPO was 250 U/ml. Eight standard concentrations were prepared from a stock 20 dilution of 20 U/ml sufficient for 4.5 ml of each standard or 20 assays in duplicate.  $20 \text{ U/ml} = 88 \mu\text{l} @ 250 \text{ U/ml} + 1012 \mu\text{l} \text{ dilution buffer}$ . The eight standard concentrations are shown in Table 2. The standards were divided into 450  $\mu\text{l}$  aliquots in microcentrifuge tubes and 25 stored at  $-20^\circ\text{C}$  until used.

Antibody 1 (1 mg) was dissolved in 1 ml RIA/DB and stored in 20  $\mu\text{l}$  aliquots in 0.5 ml microcentrifuge tubes at  $-20^\circ\text{C}$  until used (each aliquot was sufficient for 100 assay tubes). On the assay date, Ab1 was prepared at a 30 working concentration of  $2\mu\text{g/ml}$  with dilution buffer (a 1/500 dilution of the stock aliquot) to give a final assay concentration of  $0.67 \mu\text{g/ml}$  in 300  $\mu\text{l}$  when added as 100  $\mu\text{l}$  to all tubes except non-specific binding (NSB) and total counts (TC).

35 Since all standard and sample binding inhibition % were based on the maximum binding (MB), to avoid the possibility of having a poor reference duplicate, the MB

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was determined in quadruplicate.

The  $^{125}\text{I}$ -EPO was diluted with 4.5 ml of dilution buffer and divided into 25 x 0.2 ml aliquots in 500  $\mu\text{l}$  microcentrifuge tubes. A 5  $\mu\text{l}$  sample was added to 50  $\mu\text{l}$  of dilution buffer in an 11x75 mm tube and counted for 1 min. in the RIAGamma using program 2. The corrected CPM usually ranged between 4500 - 6000 CPM/ $\mu\text{l}$ . The tracer aliquots were stored, with lead shielding, at  $-20^{\circ}\text{C}$  until used.

Table 3 lists the assay setup and addition of reagents. The reagents were added and incubated in the following order: A) dilution buffer and NRS @ 1/33.3 were added to the appropriate tubes 3-8; B) EPO standards and sample dilutions were added to the appropriate tubes; C) antibody 1 was added to all tubes except 1-4 and; all tubes were vortexed and incubated at room temperature for 4h.  $^{125}\text{I}$ -EPO @ 6000 CPM/100  $\mu\text{l}$  was added to all tubes. As the tracer activity decreases over time, the maximum binding % decreases resulting in a loss of sensitivity and accuracy. Therefore, we used 6000 CPM total counts during the first week of a new batch and increased by 1000 CPM per week thereafter.

All tubes (except TC's) were vortexed and incubated overnight (18-20 hours) @  $4^{\circ}\text{C}$ . Ab 2 and NRS (1/62.5) was added to all tubes except TC's. All tubes (except TC's) were vortexed and incubated at room temperature for 2 hours. 1.5ml of 3.8% PEG 8000 (w/v in dilution buffer) was added to all tubes (except TC's) resulting in a final PEG concentration of 3%. All tubes (except TC's) were vortexed and incubated for 10 minutes at room temperature. All tubes were centrifuged (except TC's) at 1500 x g and  $4^{\circ}\text{C}$  for 20 minutes (1500 x g = 2800 RPM when using the IEC Centra-8R with the Cat.# 5737 12x75 mm tube adapters). The supernants were removed by aspiration and the tubes containing the pellets were counted in the gamma counter.

The standard curve was plotted using linear regression of logit (B/ $B_0$ ) vs log concentration where B = CPM bound and,  $B_0$  = maximum CPM bound (reference binding

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of Ab 1 without inhibitor). The regression line  $y = C_0 + C_1(x)$  where,  $C_0 = y$  intercept,  $C_1 =$  slope,  $x = \text{Log (Concentration)}$ ,  $y = \text{Log (R/(1-R))}$  and,  $R = B/B_0$ .

Interference of culture medium was tested by the  
5 addition of up to 95  $\mu\text{l}$  of a possible 100  $\mu\text{l}$  sample volume of Ham's F-12 containing 10% FCS. No effect on binding was observed. Recovery of EPO at 100, 300, and 600 mU/ml was 93.85, 100.41, and 96.33% respectively.

Screening was performed as follows. The assay was  
10 performed in one day for screening samples by using the following modifications to the above-noted method. Standards and sample dilutions were incubated with Ab1 for 1 hour at 37°C instead of 4 hours at room temperature.  $^{125}\text{I}$ -EPO was incubated with all tubes for 1 hour at 37°C  
15 instead of overnight at 4°C. Antibody 2 and NRS are incubated for 1 hour at 37°C instead of 2 hours at room temperature. The remaining procedures were as described above and the results are given below.

The first transfection series was performed with the  
20 following target vectors: PLW24 (EF1 promoter,  $\text{EPO}_{\text{long}}$ ) referred to as EPO-1 or; p24MAR-1 (EF1 promoter,  $\text{EPO}_{\text{long}}$ , 5'apoB & 3' 60 bp-deleted apoB SAR elements) referred to as EPO-1'. The selection plasmid was pSV2neo. The target:selection plasmid ratio was 5:1 (30  $\mu\text{g}$ :6 $\mu\text{g}$ ).

25 One sample was picked up randomly from each transfection and was cloned by the dilution method. EPO-1-0-1 was cloned from the EPO-1 transfection and EPO-1-0-4' from the EPO-1' transfection (' indicates the presence of SAR elements in the target vector).

30 After the first round of cloning, the two EPO-1 clones and four EPO-1' clones expressing the highest levels of EPO were selected and expanded. Levels of EPO production in the medium, determined by the RIA assays were as follows: EPO-1-1-5: 8 u/ $10^6$  cells/day; EPO-1-1-6:  
35 9 u/ $10^6$  cells/day; EPO-1-1-7': 41 u/ $10^6$  cells/day; EPO-1-1-8': 30 u/ $10^6$  cells/day; EPO-1-1-13': 125 u/ $10^6$  cells/day and; EPO-1-1-14': 81 u/ $10^6$  cells/day.



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After a second round of cloning, levels of EPO expression were as follows: EPO-1-2-22: 5 u/10<sup>6</sup> cells/day; EPO-1-2-23: 7 u/10<sup>6</sup> cells/day; EPO-1-2-26: 13 u/10<sup>6</sup> cells/day; EPO-1-2-27: 19 u/10<sup>6</sup> cells/day; EPO-1-2-15\*: 13 u/10<sup>6</sup> cells/day; EPO-1-2-16\*: 20 u/10<sup>6</sup> cells/day and; EPO-1-2-17\*: 105 u/10<sup>6</sup> cells/day.

The second transfection series was performed with the same target and selection vectors as the first transfection series. The target plasmid:selection plasmid ratio was 12:1 (30µg:2.5µg).

At day 10 after transfection, EPO-2-0-11 (producing 0.8 u/ml EPO) from EPO-2 transfection (target plasmid has no SAR elements) and EPO 2-0-9\* (producing 0.6 u/ml) from EPO-2\* transfection (target plasmid with SAR elements) were selected for cloning. 21 days after cloning, the levels of EPO production in the medium for EPO-2-0-11 were as follows: 36/48 samples had <160 u/ml; 11/48 samples had 160-800 u/ml; and 1/48 samples had 800 u/ml. Levels of EPO production for EPO-2-0-9\* were as follows: 47/48 samples had 1,000-4,000 u/ml and; 1/48 samples had <800 u/ml.

Two of the EPO-2 clones and three of the EPO-2\* clones (expressing the highest levels of EPO per cell cluster) were expanded, and the levels of expression were as follows: EPO-2-1-24 (514 u/ml 21 days after cloning) and 109 u/10<sup>6</sup> cells/day after expansion; EPO-2-1-25 (800 u/ml 21 days after cloning) and 182 u/10<sup>6</sup> cells/day after expansion; EPO-2-1-18\* (1,962 u/ml 21 days after cloning) and 185 u/10<sup>6</sup> cells/day after expansion; EPO-2-1-19\* (1,682 u/ml 21 days after cloning) and 1,306 u/10<sup>6</sup> cells/day after expansion and; EPO-2-1-21\* (1,777 u/ml 21 days after cloning) and 332 u/10<sup>6</sup> cells/day after expansion.

EPO-2-1-19\* was found to maintain its levels of EPO production while submitted to successive freezing-revival cycles as follows: 962 and 1,038 u/10<sup>6</sup> cells/day after the first cycle; 1,323 and 1,264 u/10<sup>6</sup> cells/day (between days 3 and 4) after the second cycle and; 1,188 u/10<sup>6</sup> cells/day

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(between days 3 and 4) after the third cycle.

EPO-2-1-19\* was submitted to a second round of cloning. The clones producing the highest levels of EPO produced between 1,500 and 1,700 u/10<sup>6</sup> cells/day (between  
5 days 3 and 4). The EPO-2-1-19\* cell line has remained stable and expressed constant levels of EPO for about eight months.

# TABLE 1

## Assembly of expression vectors

EF1 promoter	CMV promoter	EPO sequence	5' SAR element	3' SAR element
pLW24 Rh49	pLW25 Rh216	cDNA <sub>long</sub>	-	-
p24MAR-1 Rh163	p25MAR-1 Rh169	cDNA <sub>long</sub>	Rh10	Rh32
pAP142 Rh223	pAP140 Rh233	cDNA <sub>long</sub>	Rh32	Rh32
pAP59 Rh183	pAP67 Rh209	genomic	-	-
pAP123 Rh211	pAP127 Rh206	genomic	Rh32	Rh32
pAP132 Rh221	pAP134 Rh222	genomic	Rh10	Rh32

# SUBSTITUTE SHEET

TABLE 2

Standard No.	20 U/ml Stock ( $\mu$ l)	Dilution Buffer ( $\mu$ l)	Standard Concentration (mU/ml)
1	4.5	4495.5	20
2	9.0	4491.0	40
3	18.0	4482.0	80
4	36.0	4464.0	160
5	67.5	4432.5	300
6	135.0	4365.0	600
7	270.0	4230.0	1,200
8	450.0	4050.0	2,000

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TABLE 3

Tube #	Code	Dilution Buffer (μl)	NRS @ 1/33.3 (μl)	EPO Standard (μl)	Ab 1 @ 2μg/ml (μl)	<sup>125</sup> I-EPO 6000 CPM (μl)	Ab 2 @ 1/5 (μl)	NRS @ 1/62.5 (μl)
1-2	TC	0	0	0	0	100	0	0
3-4	NSB	100	100	0	0	100	50	50
5-8	MB	100	0	0	100	100	50	50
<b>Standard Curve</b>								
9-10	20 mU/ml	0	0	100	100	100	50	50
11-12	40 mU/ml	0	0	100	100	100	50	50
13-14	80 mU/ml	0	0	100	100	100	50	50
15-16	160 mU/ml	0	0	100	100	100	50	50
17-18	300 mU/ml	0	0	100	100	100	50	50
19-20	600 mU/ml	0	0	100	100	100	50	50
21-22	1200 mU/ml	0	0	100	100	100	50	50
23-24	2000 mU/ml	0	0	100	100	100	50	50
<b>Inter/Intra-assay Quality Controls</b>								
25-26	100 mU/ml	0	0	100	100	100	50	50
27-28	400 mU/ml	0	0	100	100	100	50	50
29-30	800 mU/ml	0	0	100	100	100	50	50
<b>Unknowns</b>								
31-32	Sample 1	100 μl of Sample Dilution	0	0	100	100	50	50

SUBSTITUTE SHEET

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## SEQUENCE LISTING

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## (v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk  
(B) COMPUTER: IBM PC compatible  
(C) OPERATING SYSTEM: PC-DOS/MS-DOS  
(D) SOFTWARE: PatentIn Release #1.0, Version #1.25

## (vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER:  
(B) FILING DATE:  
(C) CLASSIFICATION:

## (viii) ATTORNEY/AGENT INFORMATION:

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## (2) INFORMATION FOR SEQ ID NO:1:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 51 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

- 45 -

(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

(B) CLONE: EPO1

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

AGCTTGCCCG GGATGAGGGC CACCGGTGTG GTCACCCGGC GCGCCCCAGG T

51

(2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 44 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

(B) CLONE: EPO2

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

CGCTGAGGGA CCCCGGCCAG GCGCGGAGAT GGGGGTGCAC GAAT

44

(2) INFORMATION FOR SEQ ID NO:3:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 51 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

(B) CLONE: EPO3

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

GTCCTGCCTG GCTGTGGCTT CTCCTGTCCC TGCTGTCGCT CCCTCTGGGC C

51

(2) INFORMATION FOR SEQ ID NO:4:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 59 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

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(B) CLONE: EPO4

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

TCCCAGTCCT GGGCGCCCCA CCACGCCTCA TCTGTGACAG CCGAGTCCTG GAGAGGTAC 59

(2) INFORMATION FOR SEQ ID NO:5:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 56 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

(B) CLONE: EPO5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

CTCTTGAGG CCAAGGAGGC CGAGAATATC ACGACGGGCT GTGCTGAACA TTGCAG 56

(2) INFORMATION FOR SEQ ID NO:6:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 58 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

(B) CLONE: EPO6

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

CTTGAATGAG AATATCACTG TCCCAGACAC CAAAGTTAAT TTCTATGCCT GGAAGAGG 58

(2) INFORMATION FOR SEQ ID NO:7:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 55 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

(B) CLONE: EPO7



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## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

ATGGAGGTCG GGCAGCAGGC CGTAGAAGTC TGGCAGGGCC TGGCCCTGCT GTCGG

55

## (2) INFORMATION FOR SEQ ID NO:8:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 61 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA

## (vi) ORIGINAL SOURCE:

- (A) ORGANISM: Homo sapiens

## (vii) IMMEDIATE SOURCE:

- (B) CLONE: EPO8

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

AAGCTGTCCT GCGGGGCCAG GCCCTGTTGG TCAACTCTTC CCAGCCGTGG GAGCCCCTGC

60

A

61

## (2) INFORMATION FOR SEQ ID NO:9:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 54 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA

## (vi) ORIGINAL SOURCE:

- (A) ORGANISM: Homo sapiens

## (vii) IMMEDIATE SOURCE:

- (B) CLONE: EPO9

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

GCTGCATGTG GATAAAGCCG TCAGTGGCCT TCGCAGCCTC ACCACTCTGC TTCG

54

## (2) INFORMATION FOR SEQ ID NO:10:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 51 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA

## (vi) ORIGINAL SOURCE:

- (A) ORGANISM: Homo sapiens

## (vii) IMMEDIATE SOURCE:

- (B) CLONE: EPO10

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

- 48 -

GGCTCTGGGA GCCCAGAAGG AAGCCATCTC CCCTCCAGAT GCGGCCTCAG C

51

## (2) INFORMATION FOR SEQ ID NO:11:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 51 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

- (vi) ORIGINAL SOURCE:
  - (A) ORGANISM: Homo sapiens

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: EPO11

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

TGCTCCACTC CGAACAATCA CTGCTGACAC TTTCCGCAAA CTCTTCCGAG T

51

## (2) INFORMATION FOR SEQ ID NO:12:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 47 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

- (vi) ORIGINAL SOURCE:
  - (A) ORGANISM: Homo sapiens

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: EPO12

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

CTACTCCAAT TTCCTCCGGG GAAAGCTGAA GCTGTACACA GGGGAGG

47

## (2) INFORMATION FOR SEQ ID NO:13:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 48 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

- (vi) ORIGINAL SOURCE:
  - (A) ORGANISM: Homo sapiens

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: EPO13

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

CCTGCAGGAC AGGGGACAGA TGACCAGGTG TGTCGACCTG GGCATATC

48

## (2) INFORMATION FOR SEQ ID NO:14:

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- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 48 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

- (ii) MOLECULE TYPE: cDNA

- (vi) ORIGINAL SOURCE:
  - (A) ORGANISM: Homo sapiens

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: EPO13b

- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

CCTGCAGGAC AGGGGACAGA TGACCAGGTG TGTCCACCTG GGCATATC

48

(2) INFORMATION FOR SEQ ID NO:15:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 50 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

- (ii) MOLECULE TYPE: cDNA

- (vi) ORIGINAL SOURCE:
  - (A) ORGANISM: Homo sapiens

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: EPO15

- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

CACCACCTCC CTCACCAATA TTGCTTGTGC CACACCCTCC CCCGCCACTC

50

(2) INFORMATION FOR SEQ ID NO:16:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 52 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

- (ii) MOLECULE TYPE: cDNA

- (vi) ORIGINAL SOURCE:
  - (A) ORGANISM: Homo sapiens

- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: EPO15

- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

CTGAACCCCG TCGAGGGGCT CTCAGCTCAG CGCCAGCCTG TCCCATGGAC CA

52

(2) INFORMATION FOR SEQ ID NO:17:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 55 base pairs
  - (B) TYPE: nucleic acid

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(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:  
(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:  
(B) CLONE: EPO1 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

CCTCAGCGAC CTGGGGCGCG CCGGGTGACC ACACCGGTGG CCCTCATCCC GGGCA

55

(2) INFORMATION FOR SEQ ID NO:18:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 44 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:  
(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:  
(B) CLONE: EPO2 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

GGCAGGACAT TCGTGCACCC CCATCTCCGC GCCTGGCCCGG GGTC

44

(2) INFORMATION FOR SEQ ID NO:19:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 52 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:  
(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:  
(B) CLONE: EPO3 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

GGACTGGGAG GCCCAGAGGG AGCGACAGCA GGGACAGGAG AAGCCACAGC CA

52

(2) INFORMATION FOR SEQ ID NO:20:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 46 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

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- (ii) MOLECULE TYPE: cDNA
- (vi) ORIGINAL SOURCE:
  - (A) ORGANISM: Homo sapiens
- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: EPO4 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:

CTCTCCAGGA CTCGGCTGTC ACAGATGAGG CGTGGTGGGG CGCCCA

46

(2) INFORMATION FOR SEQ ID NO:21:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 52 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (vi) ORIGINAL SOURCE:
  - (A) ORGANISM: Homo sapiens
- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: EPO5 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

TTCAGCACAG CCCGTCGTGA TATTCTCGGC CTCCTTGGCC TCCAAGAGGT AC

52

(2) INFORMATION FOR SEQ ID NO:22:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 58 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (vi) ORIGINAL SOURCE:
  - (A) ORGANISM: Homo sapiens
- (vii) IMMEDIATE SOURCE:
  - (B) CLONE: EPO6A

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

AGGCATAGAA ATTAACCTTG GTGTCTGGGA CAGTGATATT CTCATTCAAG CTGCAATG

58

(2) INFORMATION FOR SEQ ID NO:23:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 55 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (vi) ORIGINAL SOURCE:

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(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:  
(B) CLONE: EPO7 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

AGGGCCAGGC CCTGCCAGAC TTCTACGGCC TGCTGCCCCGA CCTCCATCCT CTTCC 55

(2) INFORMATION FOR SEQ ID NO:24:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 65 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:  
(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:  
(B) CLONE: EPO8 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

GGGGCTCCCA CGGCTGGGAA GAGTTGACCA ACAGGGCCTG GCCCCGCAGG ACAGCTTCCG 60  
ACAGC 65

(2) INFORMATION FOR SEQ ID NO:25:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 50 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:  
(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:  
(B) CLONE: EPO9 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

AGTGGTGAGG CTGCGAAGGC CACTGACGGC TTTATCCACA TGCAGCTGCA 50

(2) INFORMATION FOR SEQ ID NO:26:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 51 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:  
(A) ORGANISM: Homo sapiens

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(vii) IMMEDIATE SOURCE:  
(B) CLONE: EPO10 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

CGCATCTGGA GGGGAGATGG CTTCTTCTG GGCTCCCAGA GCGCGAAGCA G

51

(2) INFORMATION FOR SEQ ID NO:27:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 61 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:  
(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:  
(B) CLONE: EPO11 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:

AGACTCGGAA GAGTTTGCGG AAAGTGTGAG CAGTGATTGT TCGGAGTGGG GCAGCTGAGG 60  
C 61

(2) INFORMATION FOR SEQ ID NO:28:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 53 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:  
(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:  
(B) CLONE: EPO 12 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:

CCTGCAGGCC TCCCCTGTGT ACAGCTTCAG CTTTCCCCGG AGGAAATTGG AGT

53

(2) INFORMATION FOR SEQ ID NO:29:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 47 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:  
(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

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(B) CLONE: EPO13 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:

GGTGGTGGAT ATGCCCAGGT CGACACACCT GGTCATCTGT CCCCTGT

47

(2) INFORMATION FOR SEQ ID NO:30:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 47 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

(B) CLONE: EPO13b alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

GGTGGTGGAT ATGCCCAGGT GGACACACCT GGTCATCTGT CCCCTGT

47

(2) INFORMATION FOR SEQ ID NO:31:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 51 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

(B) CLONE: EPO14 alpha

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

GGGTTTCAGGA GTGGCGGGGG AGGGTGTGGC ACAAGCAATA TTGGTGAGGG A

51

(2) INFORMATION FOR SEQ ID NO:32:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 48 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

(B) CLONE: EPO15 alpha



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## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:

AGCTTGGTCC ATGGGACAGG CTGGCGCTGA GCTGAGAGCC CCTCGACG

48

## (2) INFORMATION FOR SEQ ID NO:33:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 3602 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: DNA (genomic)

## (vi) ORIGINAL SOURCE:

- (A) ORGANISM: Homo sapiens

## (ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: join(625..637, 1201..1346, 1605..1691, 2303..2482, 2617..2772)

## (ix) FEATURE:

- (A) NAME/KEY: mRNA
- (B) LOCATION: join(625..637, 1201..1346, 1605..1691, 2303..2482, 2617..2772)

## (ix) FEATURE:

- (A) NAME/KEY: mat\_peptide
- (B) LOCATION: join(1269..1346, 1605..1691, 2303..2482, 2617..2769)

## (ix) FEATURE:

- (A) NAME/KEY: prim\_transcript
- (B) LOCATION: join(625..3337)

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:

```

AAGCTTCTGG GCTTCCAGAC CCAGCTACTT TGCGGAAGTC AGCAACCCAG GCATCTCTGA   60
GTCTCCGCCC AAGACCGGGA TGCCCCCAG GGGAGGTGTC CGGGAGCCCA GCCTTTCCCA  120
GATAGCACGC TCCGCCAGTC CCAAGGGTGC GCAACCGGCT GCACTCCCCT CCCGCGACCC  180
AGGGCCCCGGG AGCAGCCCC ATGACCCACA CGCACGTCTG CAGCAGCCCC GCTCACGCCC  240
CGGCGAGCCT CAACCCAGGC GTCCTGCCCC TGCTCTGACC CCGGGTGGCC CCTACCCCTG  300
GCGACCCCTC ACGCACACAG CCTCTCCCC ACCCCCACCC GCGCACGCAC ACATGCAGAT  360
AACAGCCCCG ACCCCCGGCC AGAGCCGCAG AGTCCCTGGG CCACCCCGGC CGCTCGCTGC  420
GCTGCGCCGC ACCGCGCTGT CCTCCCGGAG CCGGACCGGG GCCACCGCGC CCGCTCTGCT  480
CCGACACCGC GCCCCCTGGA CAGCCGCCCT CTCCTCTAGG CCCGTGGGGC TGGCCCTGCA  540
CCGCCGAGCT TCCCGGGATG AGGGCCCCCG GTGTGGTCAC CCGGCGCGCC CCAGGTCGCT  600
GAGGGACCCC GGCCAGGCGC GGAG ATG GGG GTG CAC  G GTGAGTACTC           647
                               Met Gly Val His
                               -27   -25
GCGGGCTGGG CGCTCCCGCC GCCCGGTTC CTGTTTGGAG GGGGATTTAG CGCCCCGGCT  707

```

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BNSDOCID: <WO 9619573A1 | >

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TGCATACCTT CTGTTTGCTC AGCTTGGTGC TTGGGGCTGC TGAGGGGCAG GAGGGAGAGG 2261  
 GTGACATGGG TCAGCTGACT CCCAGAGTCC ACTCCCTGTA G GTC GGG CAG CAG 2314  
 Val Gly Gln Gln  
 GCC GTA GAA GTC TGG CAG GGC CTG GCC CTG CTG TCG GAA GCT GTC CTG 2362  
 Ala Val Glu Val Trp Gln Gly Leu Ala Leu Leu Ser Glu Ala Val Leu  
 60 65 70 75  
 CGG GGC CAG GCC CTG TTG GTC AAC TCT TCC CAG CCG TGG GAG CCC CTG 2410  
 Arg Gly Gln Ala Leu Val Asn Ser Ser Gln Pro Trp Glu Pro Leu  
 80 85 90  
 CAG CTG CAT GTG GAT AAA GCC GTC AGT GGC CTT CGC AGC CTC ACC ACT 2458  
 Gln Leu His Val Asp Lys Ala Val Ser Gly Leu Arg Ser Leu Thr Thr  
 95 100 105  
 CTG CTT CGG GCT CTG GGA GCC CAG GTGAGTAGGA GCGGACACTT CTGCTTGCCC 2512  
 Leu Leu Arg Ala Leu Gly Ala Gln  
 110 115  
 TTTCTGTAAG AAGGGGAGAA GGGTCTTGCT AAGGAGTACA GGAAGTGTCC GTATTCCTTC 2572  
 CCTTTCTGTG GCACTGCAGC GACCTCCTGT TTTCTCCTTG GCAG AAG GAA GCC ATC 2628  
 Lys Glu Ala Ile  
 TCC CCT CCA GAT GCG GCC TCA GCT GCT CCA CTC CGA ACA ATC ACT GCT 2676  
 Ser Pro Pro Asp Ala Ala Ser Ala Ala Pro Leu Arg Thr Ile Thr Ala  
 120 125 130 135  
 GAC ACT TTC CGC AAA CTC TTC CGA GTC TAC TCC AAT TTC CTC CGG GGA 2724  
 Asp Thr Phe Arg Lys Leu Phe Arg Val Tyr Ser Asn Phe Leu Arg Gly  
 140 145 150  
 AAG CTG AAG CTG TAC ACA GGG GAG GCC TGC AGG ACA GGG GAC AGA TGACCAGGTG 2779  
 Lys Leu Lys Leu Tyr Thr Gly Glu Ala Cys Arg Thr Gly Asp Arg  
 155 160 165  
 TGTCCACCTG GGCATATCCA CCACCTCCCT CACCAACATT GCTTGTGCCA CACCCTCCCC 2839  
 CGCCACTCCT GAACCCCGTC GAGGGGCTCT CAGCTCAGCG CCAGCCTGTC CCATGGACAC 2899  
 TCCAGTGCCA GCAATGACAT CTCAGGGGCC AGAGGAACTG TCCAGAGAGC AACTCTGAGA 2959  
 TCTAAGGATG TCACAGGGCC AACTTGAGGG CCCAGAGCAG GAAGCATTCA GAGAGCAGCT 3019  
 TTAAACTCAG GGACAGAGCC ATGCTGGGAA GACGCCTGAG CTCACTCGGC ACCCTGCAAA 3079  
 ATTTGATGCC AGGACACGCT TTGGAGGCGA TTTACCTGTT TTCGCACCTA CCATCAGGGA 3139  
 CAGGATGACC TGGAGAACTT AGGTGGCAAG CTGTGACTTC TCCAGGTCTC ACGGGCATGG 3199  
 GCACTCCCTT GGTGGCAAGA GCCCCTTGA CACCGGGGTG GTGGGAACCA TGAAGACAGG 3259  
 ATGGGGGCTG GCCTCTGGCT CTCATGGGGT CCAAGTTTTG TGTATTCTTC AACCTCATTG 3319  
 ACAAGAACTG AAACCACCAA TATGACTCTT GGCTTTTCTG TTTTCTGGGA ACCTCCAAAT 3379  
 CCCCTGGCTC TGTCCCACTC CTGGCAGCAG TGCAGCAGGT CCAGGTCCGG GAAATGAGGG 3439  
 GTGGAGGGGG CTGGGCCCTA CGTGCTGTCT CACACAGCCT GTCTGACCTC TCGACCTACC 3499

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GGCCTAGGCC ACAAGCTCTG CCTACGCTGG TCAATAAGGT GTCTCCATTC AAGGCCTCAC 3559  
 CGCAGTAAGG CAGCTGCCAA CCCTGCCCAG GGCAAGGCTG CAG 3602

## (2) INFORMATION FOR SEQ ID NO:34:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 193 amino acids  
 (B) TYPE: amino acid  
 (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:

Met Gly Val His Glu Cys Pro Ala Trp Leu Trp Leu Leu Leu Ser Leu  
 -27 -25 -20 -15  
 Leu Ser Leu Pro Leu Gly Leu Pro Val Leu Gly Ala Pro Pro Arg Leu  
 -10 -5 1 5  
 Ile Cys Asp Ser Arg Val Leu Glu Arg Tyr Leu Leu Glu Ala Lys Glu  
 10 15 20  
 Ala Glu Asn Ile Thr Thr Gly Cys Ala Glu His Cys Ser Leu Asn Glu  
 25 30 35  
 Asn Ile Thr Val Pro Asp Thr Lys Val Asn Phe Tyr Ala Trp Lys Arg  
 40 45 50  
 Met Glu Val Gly Gln Gln Ala Val Glu Val Trp Gln Gly Leu Ala Leu  
 55 60 65  
 Leu Ser Glu Ala Val Leu Arg Gly Gln Ala Leu Leu Val Asn Ser Ser  
 70 75 80 85  
 Gln Pro Trp Glu Pro Leu Gln Leu His Val Asp Lys Ala Val Ser Gly  
 90 95 100  
 Leu Arg Ser Leu Thr Thr Leu Leu Arg Ala Leu Gly Ala Gln Lys Glu  
 105 110 115  
 Ala Ile Ser Pro Pro Asp Ala Ala Ser Ala Ala Pro Leu Arg Thr Ile  
 120 125 130  
 Thr Ala Asp Thr Phe Arg Lys Leu Phe Arg Val Tyr Ser Asn Phe Leu  
 135 140 145  
 Arg Gly Lys Leu Lys Leu Tyr Thr Gly Glu Ala Cys Arg Thr Gly Asp  
 150 155 160 165  
 Arg

## (2) INFORMATION FOR SEQ ID NO:35:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 788 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA

## (vi) ORIGINAL SOURCE:

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(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

(B) CLONE: EP0long

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

```
AGCTTGCCCG GGATGAGGGC CACCGGTGTG GTCACCCGGC GCGCCCCAGG TCGCTGAGGG 60
ACCCCGGCCA GGC GCGGAGA TGGGGGTGCA CGAATGTCCT GCCTGGCTGT GGCTTCTCCT 120
GTCCCTGCTG TCGCTCCCTC TGGGCCTCCC AGTCTGGGC GCCCACCAC GCCTCATCTG 180
TGACAGCCGA GTCCTGGAGA GGTACCTCTT GGAGGCCAAG GAGGCCGAGA ATATCACGAC 240
GGGCTGTGCT GAACATTGCA GCTTGAATGA GAATATCACT GTCCCAGACA CCAAAGTTAA 300
TTTCTATGCC TGGAAGAGGA TGGAGGTCGG GCAGCAGGCC GTAGAAGTCT GGCAGGGCCT 360
GGCCCTGCTG TCGGAAGCTG TCCTGCGGGG CCAGGCCCTG TTGGTCAACT CTTCCCAGCC 420
GTGGGAGCCC CTGCAGCTGC ATGTGGATAA AGCCGTCAGT GGCCTTCGCA GCCTCACCAC 480
TCTGCTTCGG GCTCTGGGAG CCCAGAAGGA AGCCATCTCC CCTCCAGATG CGGCCTCAGC 540
TGCTCCACTC CGAACAATCA CTGCTGACAC TTTCCGAAA CTCTTCCGAG TCTACTCCAA 600
TTTCTCCGG GGAAAGCTGA AGCTGTACAC AGGGGAGGCC TGCAGGACAG GGGACAGATG 660
ACCAGGTGTG TCCACCTGGG CATATCCACC ACCTCCCTCA CCAATATTGC TTGTGCCACA 720
CCCTCCCCCG CCACTCCTGA ACCCCGTCGA GGGGCTCTCA GCTCAGCGCC AGCCTGTCCC 780
ATGGACCA 788
```

(2) INFORMATION FOR SEQ ID NO:36:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 605 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:

(B) CLONE: Rh 32

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:

```
AAAAAGATGA GGTAATTGTG TTTTATAAT TAAATATTTT ATAATTAAAA TATTTATAAT 60
TAAATATTTT ATAATTAAAT ATTTTATAAT TAAATATTTT ATAATTAAAT ATTTTATAAT 120
TAAATATTTT ATAATTAAAT ATTTTATAAT TAAATATTTT ATAATTAAAT ATTTTATAAT 180
TAAATATTTT ATAATTAAAT ATTTTATAAT TAAATATTTT ATAATTAAAT ATTTTATAAT 240
TAAATATTTT ATAATTAAAT ATTTTATAAT TAAATGTTT ATAATTACAT ATTTTATAAT 300
TAAATGTTT ATAATTACAT ATTTTATAAT TAAATGTTT ATAATTACAT ATTTTATAAT 360
```

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TAAAATGTTT ATAATTACAT ATTTTATAAT TACATATTTT ATAAAGTATT TATAATTACA 420  
TATTTTATAA TTAAAGTATT TATAATTACA TATTTTATAA TTAAAGTATT TATAATTACA 480  
TATTTTATAA TTCAATATTT TATAAATAGT TAAAAAGACG AGGAAAAAAT TAAAAAGACG 540  
AGGTTATTGA TCTCAGGAAT TGTATTTGCC AAGTGAGAAG GAAAAAATAT TCACAAAGGC 600  
TTGTA 605

## (2) INFORMATION FOR SEQ ID NO:37:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 2529 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:  
(A) ORGANISM: Homo sapiens

(vii) IMMEDIATE SOURCE:  
(B) CLONE: Rh 10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

TCTAGACCCC AGTTTCTCTA TAAGATGAGA ATATTAGTCA CGATTTGGTT TCTAAGATCC 60  
TGTCTATGTT TGAGACTACA GATACCTGTT GCTACATTTC CCTTCATAGC TCTGAACAAG 120  
GAGAATTCAG CCAATTCTC ATGGCCTTCT AAACAATCCA GAGTTTCAGT GCCATAAGGT 180  
ACTACAATTT AGTGTCAAAT TAAGTCAAAG GCTTCATTAG CCTGAAAGCT CTGTCCCTGG 240  
CCTGGGCATG GCAAACGTGA TCCCCCACTG ACCATCCCCC TGTCTCCCTT CTCCCCAGAG 300  
ACTCCAGTAG CCTGGCGTCA TCACAGGGGC CAGACATATC CAACATGTTC CCAGCTTCCT 360  
GCCACTTGAC TTTCAGTGTG CCTCCCTCTT CAGTTACCCA AATCCTGCCC ACCATTCCAG 420  
AGCCAGTTCA ATCTCACCCA TCCAGGACCC CCGAGACCCC CATCGTACCA CTATAGTCTA 480  
ACTGTGGTGT AGACCCCACTA CTGGGCACAT TGCGTACGCT CATTATTGGC TGTGACGTCT 540  
GATTATGCCC TTCTCCTGGT CTGGAAGCTC TCGGAGGTGC TCCATAATAC ATGAAGAGAA 600  
GTAGTGCTGG TGTGGGAATA GTGAGGTGTG TTTATCCATC CAGCTATCCG GCACCAGCAC 660  
TGGTCTCAGC TTTCTGAGGT AACACGTTCT GAGCCTTAGT CTTGAGAGAA CATAAAGAAA 720  
ACTTTTTTTA AAAGTAGTAA AAAGTGGCTG ACAAAGCTG ACCAAAAGCC TTCAAAGAA 780  
ATGCTAAGTT ATATCTAAGA AAGTTTACCC AAGGTCAGGC AAATATGAAA CCTAAAGCTA 840  
GACGTGGGGA AGAACTTCCG GAGAGTTGCA ATTCCCTGTG CCCCAGCATC CCCAGGAGGG 900  
CATGCCCACA TCTGATTTAG AAATCTGTGT AAAATGAGTG AAGGTTTCTA TTTCTTGGGC 960  
AGTGTGGGCA CAGGTCTTTG GAGAGGTGCA TGGCCTCCCA TAAAATCCTT CCTGCTTGAT 1020  
GGTTCCTGGAT CCTCAGCCAC AGCTCCTAAT AGCCATGAGG TTTGAGCCCA AAATAATTTA 1080  
TGTGTTTGTT TTTTCAGCCC CAAAATTTCC ATAGAATCAA AGTAGTCAGA GCTGAATGGG 1140

- 61 -

GCTAAGAGAC CGTCCATTCC TGTCTTCTCA TCACAGATGA GGGACTGCCA CCCAGAGCCG 1200  
TAGAAACTGT CCCATGGCCC CAGTTCCCAG ACCCTTCCTC TCTCCTACAG CTCCAAGTTC 1260  
ACTGTGCATT CTAAATGAAG ATGTAAACAT AGGCAGCAAC ACTCAAGAGT AAAAATGAAG 1320  
TGTGCATATG AAAGAAACCT ATTCACATGG ACCATATTAC ATTATAATCA CAGTGTTTAC 1380  
TGCTTGAATA CCATCTGCCT GGGCTAGCAA GGGTGTCACT GAGGAAGAGA GGACAAGGGG 1440  
TACCAATCTG TGAACACAC ATGGTTCTTG CTCTCCAGC TTCTCTCTCC CATTGGCAAG 1500  
GCAACAGGTA AACACATGAA AAATCAAATA ATGCTATAAG AGAAAAATGT ATTCAGGACA 1560  
ACAACAGGTT TGTATGAAGG CCTTTCATCA TCGTTGTCCT ACCTAGAAAC TGAATGACAG 1620  
GGAATCAGAG TCACAAGCTA TGAAGTCTAA CTGGGCTGGT CCCAGAGAAA GATTCACTGC 1680  
AGTAGGTGGG GCTGCAGCCA GCCCTGGGTG GGTGGAAGGA TGACATCCAC ATAGGCAAGA 1740  
GGGTGATAAT TCACTTGCGC AGCTCCTCAC TGCACATTGA ACCCTGCTGA CTTCTGGCTT 1800  
CTCTCCCGGG AGGAACTGCG ACTCAACATT CTGACCTTAT CTCTTGGGTA GCAGAATGAT 1860  
GGAGAAGGAA AGTTTCTTTT TGCTTCTCGC AGGGGTTAAT CATCCATCTG GAATGCCTAC 1920  
ATTTGGTTGA CAATGGCTCA CCCTATCATC TTCCTCCTGA ACCATTCACC TAAATGTGCC 1980  
ATTTCTTTCC TGATAGTTCT CATTTGTGTG TGTGTGTGTG TGTGTGTGTG TGCACGTGCT 2040  
CACACATGCA TGCTGTCACT GGGTAAACAG GCCACCCTGG GCACAGTTCC ATCTACAATG 2100  
TTTGAAGTTT ACTTTCCAGC TTCTGGGCAT CATTTGCAAT TATAATGCTG TCACAGGCAG 2160  
AAACGAGATA GGCTAATTAA TCGTTGTCAA TACTGATCCC TATTTGCCAG ATGAGATTTT 2220  
GGAGCAGCAT GGCTGGGAAT AATTGGTATA GACTGTATTT CCTTGCTTTA TGTCAGTGA 2280  
AATATTTATT TAAGCATCAC GGTCGCTATG CATAAATATC CTGGAAAATG GGGTATAGCT 2340  
GAATGGTGCA GATTCATTCA TTCATATTCA GCAAATTATG TTCTAAGCAC CTAATTCACT 2400  
ACGTGAACAG CACTAACTC AGAATATTGG TCTGCTGGGG TCCTTTATTA GCTTCCATGA 2460  
TTCCCTGAAC TTGGCCAAGA CCCTTCTGGT CGGCTGCAGA TAGGCACAAT GGATAGTTTT 2520  
GCTTCTAGA 2529

I CLAIM

1. A recombinant DNA molecule adapted for transfection of a host cell comprising a nucleic acid molecule encoding mammalian erythropoietin, an expression control sequence  
5 operatively linked thereto and at least one SAR element.
2. A recombinant DNA molecule as claimed in claim 1 wherein the nucleic acid molecule encodes human erythropoietin.
3. A recombinant DNA molecule as claimed in claim 2  
10 wherein the nucleic acid molecule encodes erythropoietin having the amino acid sequence shown in SEQ ID NO 34.
4. A recombinant DNA molecule as claimed in claim 3 wherein the nucleic acid molecule has the sequence as shown in SEQ ID NO. 35.
- 15 5. A recombinant DNA molecule as claimed in claim 1 wherein the SAR element is a SAR element which co-maps with a chromatin domain boundary.
6. A recombinant DNA molecule as claimed in claim 5 wherein the SAR element is a human apolipoprotein SAR  
20 element.
7. A recombinant DNA molecule as claimed in claim 5 wherein the SAR element comprises the sequence as shown in SEQ ID NO. 36 or 37.
8. A recombinant DNA molecule as claimed in claim 1  
25 wherein the nucleic acid molecule encoding mammalian erythropoietin and the expression control sequence operatively linked thereto are flanked by SAR elements.
9. An expression vector comprising a recombinant DNA molecule as claimed in claim 1.



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10. An expression vector comprising a nucleic acid molecule encoding erythropoietin and having the sequence shown in SEQ ID NO. 35 under the control of a promoter, flanked by 5' and 3' apolipoprotein SAR elements.
- 5 11. An expression vector as claimed in claim 10 wherein the promoter is a human cytomegalovirus IE enhancer and promoter, or an elongation factor-1 alpha promoter.
12. A mammalian cell transformed with an expression vector as claimed in claim 9.
- 10 13. A mammalian cell lacking a selectable gene capable of being amplified, said cell being capable of expressing recombinant EPO in vitro at levels of at least 2,000 u/10<sup>6</sup> cells in 24 hours.
14. A method of expressing recombinant mammalian  
15 erythropoietin comprising the steps of culturing a mammalian cell as claimed in claim 13 in a suitable medium until sufficient amounts of erythropoietin are produced by the cell and separating the erythropoietin produced.
15. A method of preparing recombinant erythropoietin  
20 comprising transfecting a mammalian cell with an expression vector comprising a recombinant DNA molecule as claimed in claim 1, and culturing the transfected cell in a suitable medium until sufficient amounts of erythropoietin are produced by the cell and separating the  
25 erythropoietin produced.
16. A method as claimed in claim 14 wherein the mammalian cell is further transfected with a selectable marker gene and wherein transfected cells are selected by means of the selectable marker gene.
- 30 17. A method as claimed in claim 16 wherein the

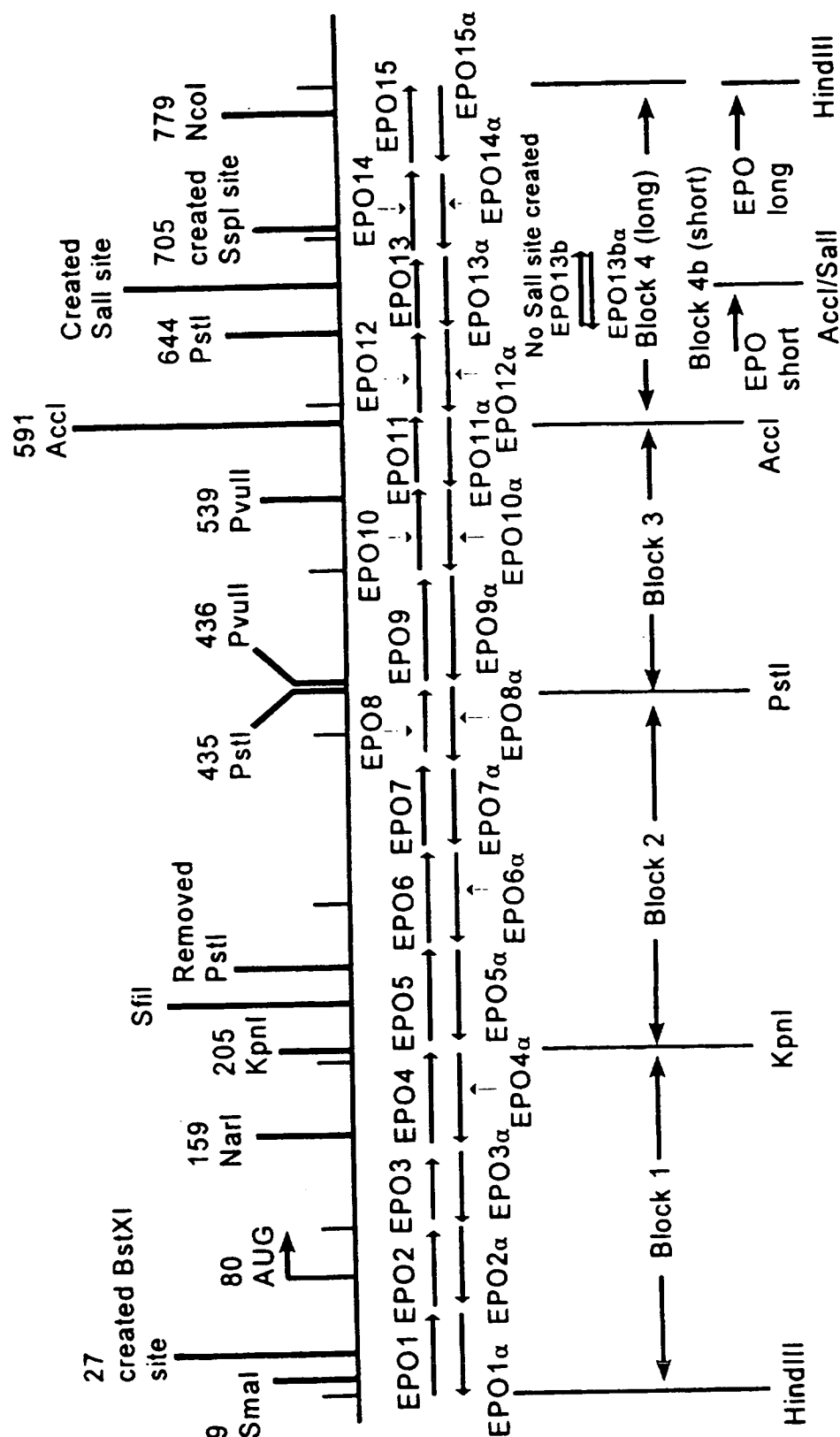
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selectable marker gene is neo.

18. A method as claimed in claim 14 comprising the additional step of identifying and selecting cells producing high levels of erythropoietin; subcloning the  
5 selected cells; establishing long term cell lines from the selected cells and; culturing the selected cells in a suitable medium until sufficient amounts of erythropoietin are produced by the cell and separating the erythropoietin produced.
- 10 19. A method as claimed in claim 14 wherein erythropoietin is produced at levels of at least 1,500 u/10<sup>6</sup> cells in 24 hours in the absence of gene amplification.
20. Erythropoietin produced by the method of claim 14.
- 15 21. A transgenic non-human animal or embryo whose germ cells and somatic cells contain a DNA construct comprising the recombinant DNA molecule as claimed in claim 1.

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FIGURE 1A



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## FIGURE 1B

OLIGONUCLEOTIDE	SEQUENCE	SEQUENCE POSITION (as per AMGEN patent cDNA)
EPO1	5'- AGC TTG CCC GGG ATG AGG GCC ACC GGT GTG GTC ACC CGG CGC GCC CCA GGT	111 - 153
EPO2	5'- CGC TGA GGG ACC CCG GCC AGG CGC GGA GAT GGG GGT GCA CGA AT	154 - 197
EPO3	5'- GTC CTG CCT GGC TGT GGC TTC TCC TGT CCC TGC TGT CGC TCC CTC TGG GCC	198 - 248
EPO4	5'- TCC CAG TCC TGG GCG CCC CAC CAC GCC TCA TCT GTG ACA GCC GAG TCC TGG AGA GGT AC	249 - 307
EPO5	5'- CTC TTG GAG GCC AAG GAG GCC GAG AAT ATC ACG ACG GGC TGT GCT GAA CAT TGC AG	308 - 363
EPO6	5'- CTT GAA TGA GAA TAT CAC TGT CCC AGA CAC CAA AGT TAA TTT CTA TGC CTG GAA GAG G	364 - 421
EPO7	5'- ATG GAG GTC GGG CAG CAG GCC GTA GAA GTC TGG CAG GGC CTG GCC CTG CTG TCG G	422 - 476

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## FIGURE 1B (Cont'd)

EPO8	5'-AAG CTG TCCTGC GGG GCC AGG CCC TGT TGG TCA ACT CTT CCC AGC CGT GGG AGC CCC TGC A	477 - 537
EPO9	5'-GCT GCA TGT GGA TAA AGC CGT CAG TGG CCT TCG CAG CCT CAC CAC TCT GCT TCG	538 - 591
EPO10	5'-GGCTCT GGG AGC CCA GAA GGA AGC CAT CTC CCC TCC AGA TGC GGC CTC AGC	592 - 642
EPO11	5'-TGC TCC ACT CCG AAC AAT CAC TGC TGA CAC TTT CCG CAA ACT CTT CCG AGT	643 - 693
EPO12	5'-CTA CTC CAA TTT CCT CCG GGG AAA GCT GAA GCT GTA CAC AGG GGA GG	694 - 740
EPO13	5'-CCT GCA GGA CAG GGG ACA GAT GAC CAG GTG TGT CGA CCT GGG CAT ATC	741 - 788
EPO13b	5'-CCT GCA GGA CAG GGG ACA GAT GAC CAG GTG TGT CCA CCT GGG CAT ATC	741 - 788
EPO14	5'-CAC CAC CTC CCT CAC CAA TAT TGC TTG TGC CAC ACC CTC CCC CGC CAC TC	789 - 838

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## FIGURE 1B (Cont'd)

EPO15	5'- CTG AAC CCC GTC GAG GGG CTC TCA GCT CAG CGC CAG CCT GTC CCA TGG ACC A	839 - 888
EPO1 $\alpha$	5'- CCT CAG CGA CCT GGG GCG CGC CGG GTG ACC ACA CCG GTG GCC CTC ATC CCG GGC A	111 - 161
EPO2 $\alpha$	5'- GGC AGG ACA TTC GTG CAC CCC CAT CTC CGC GCC TGG CCG GGG TC	162 - 205
EPO3 $\alpha$	5'- GGA CTG GGA GGC CCA GAG GGA GCG ACA GCA GGG ACA GGA GAA GCC ACA GCC A	206 - 257
EPO4 $\alpha$	5'- CTC TCC AGG ACT CGG CTG TCA CAG ATG AGG CGT GGT GGG GCG CCC A	258 - 303
EPO5 $\alpha$	5'- TTC AGC ACA GCC CGT CGT GAT ATT CTC GGC CTC CTT GGC CTC CAA GAG GTA C	304 - 355
EPO6 $\alpha$	5'- AGG CAT AGA AAT TAA CTT TGG TGT CTG GGA CAG TGA TAT TCT CAT TCA AGC TGC AAT G	356 - 413
EPO7 $\alpha$	5'- AGG GCC AGG CCC TGC CAG ACT TCT ACG GCC TGC TGC CCG ACC TCC ATC CTC TTC C	414 - 468

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## FIGURE 1B (Cont'd)

EPO8 $\alpha$	5'-GGG GCT CCC ACG GCT GGG AAG ACT TGA CCA ACA GGG CCT GGC CCC GCA GGA CAG CTT CCG ACA GC	469 - 533
EPO9 $\alpha$	5'-AGT GGT GAG GCT GCG AAG GCC ACT GAC GGC TTT ATC CAC ATG CAG CTG CA	534 - 583
EPO10 $\alpha$	5'-CGC ATC TGG AGG GGA GAT GGC TTC CTT CTG GGC TCC CAG AGC CCG AAG CAG	584 - 634
EPO11 $\alpha$	5'-AGA CTC GGA AGA GTT TCC GGA AAG TGT CAG CAG TGA TTG TTC GGA CTG GAG CAG CTG AGG C	635 - 695
EPO12 $\alpha$	5'-CCT GCA GGC CTC CCC TGT GTA CAG CTT CAG CTT TCC CCG GAG GAA ATT GGA GT	696 - 748
EPO13 $\alpha$	5'-GGT GGT GGA TAT GCC CAG GTC GAC ACA CCT GGT CAT CTG TCC CCT GT	749 - 795
EPO13b $\alpha$	5'-GGT GGT GGA TAT GCC CAG CTG GAC ACA CCT GGT CAT CTG TCC CCT GT	749 - 795
EPO14 $\alpha$	5'-GGG TTC AGG AGT GGC GGG GGA GGG TGT GGC ACA AGC AAT ATT GGT GAG GGA	796 - 846

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847 - 888

## FIGURE 1B (Cont'd)

5'-AGC TTG GTC CAT GGG ACA GGC TGG CGC  
TGA GCT GAG AGC CCC TCG ACG

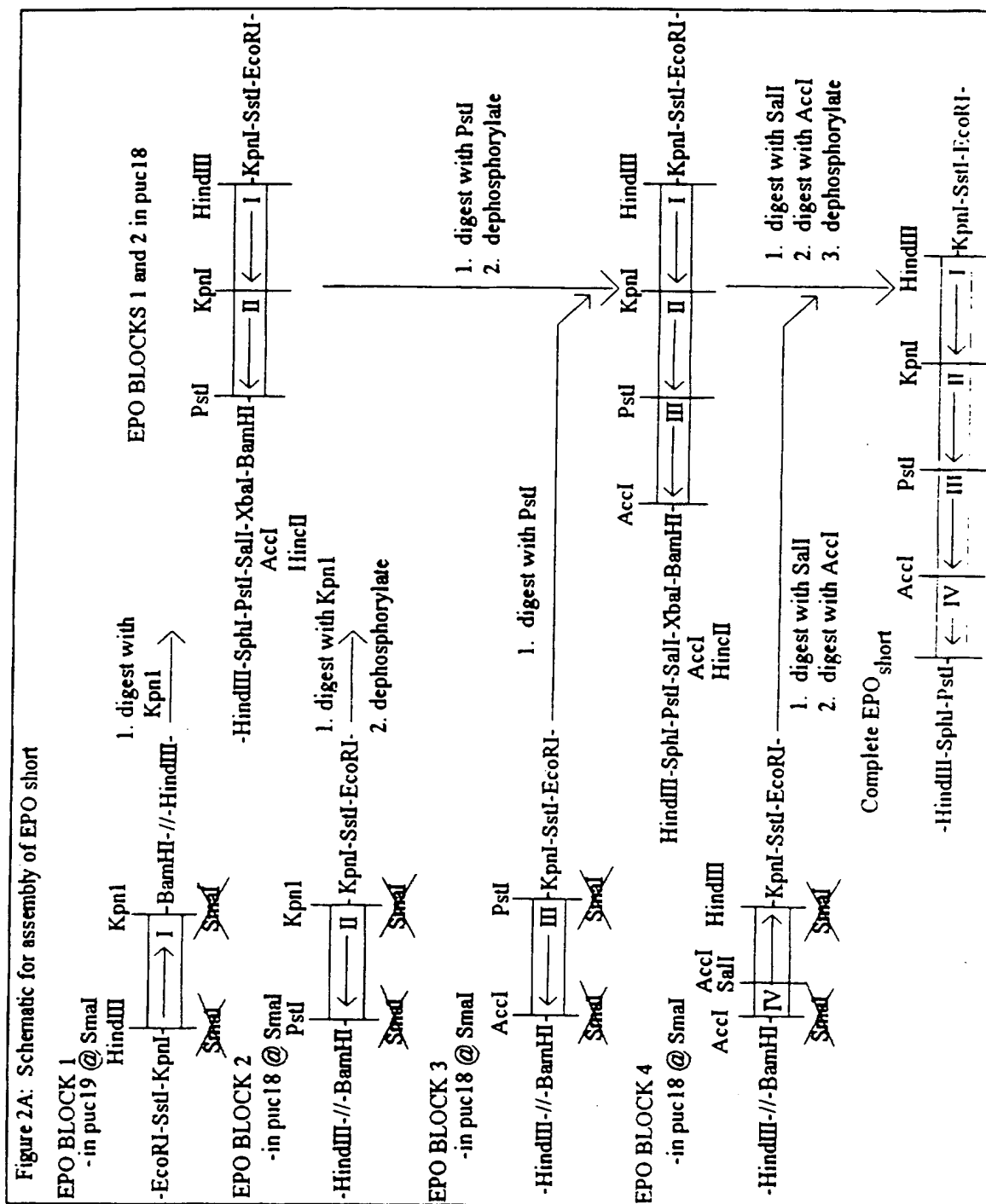
EPO15 $\alpha$

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## FIGURE 2A



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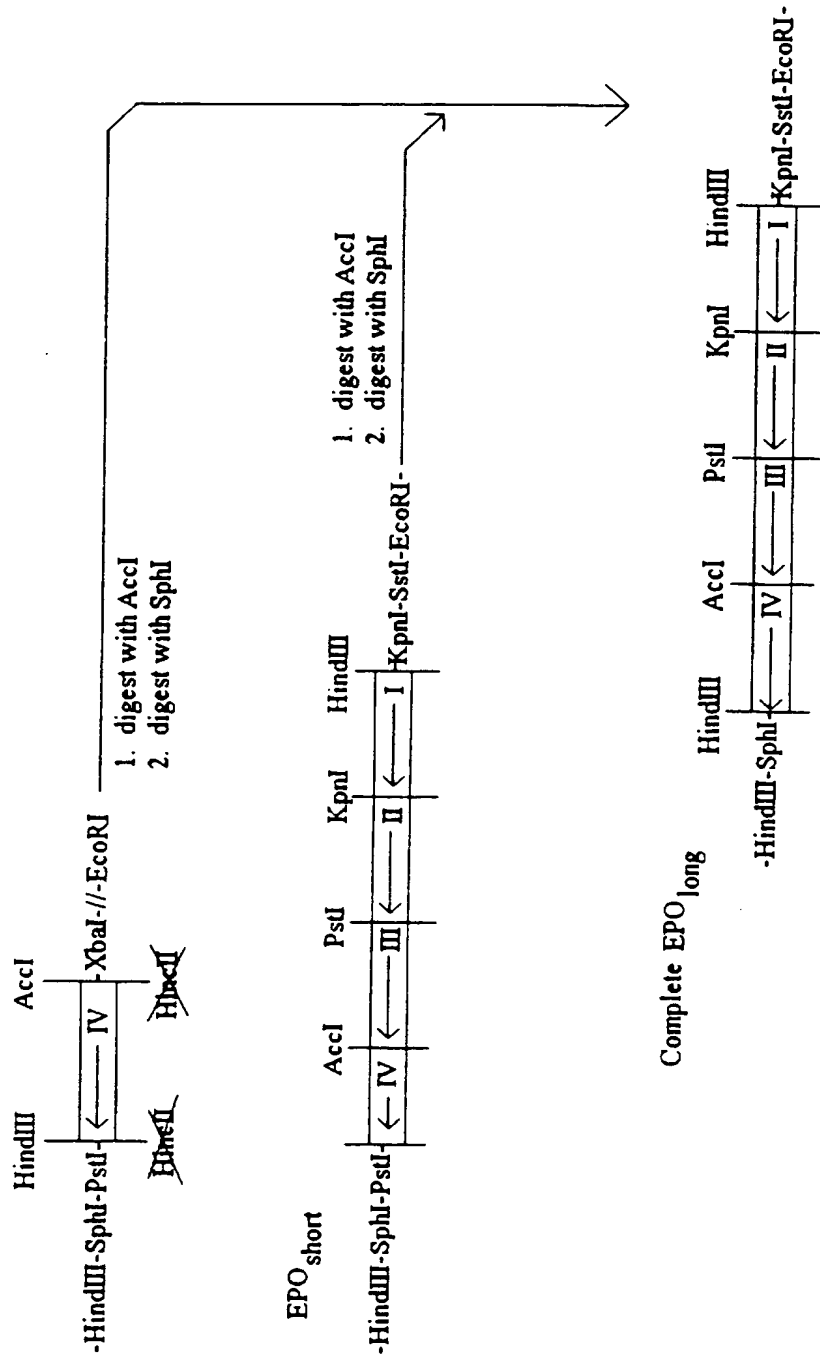
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## FIGURE 2B

Figure 2B: Schematic for assembly of EPO<sub>long</sub>

EPO BLOCK 4b

-in puc18 @ HincII



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## FIGURE 3A

1 aagctttctg gcttccagac ccagctactt tgcggaactc agcaacccag gcatctctga  
61 gtctccgccc aagaccggga tgcacccccc gggaggtgtc cgggagccca gcctttccca  
121 gatagcagc tccgccagtc ccaaggggtg gcaacccggt gcaactccct cccgcgaccc  
181 agggcccggg agcagccccc atgacccaca cgcacgtctg cagcagcccc gctcacgccc  
241 cggcgagcct caacccaggc gtccctgcccc tgcctctgacc ccgggtggcc cctacccctg  
301 ggcacccctc acgcacacag cctctccccc acccccaccc gcgcacgcac acatgcagat  
361 aacagccccg acccccggcc agagccgcag agtccctggg ccaccccggc cgctcgctgc  
421 gctgcgcccg accgcgctgt cctcccggag ccggaccggg gccaccgcgc ccgctctgct  
481 ccgacaccgc gccccctgga cagccgcccc ctccctctagg cccgtggggc tggccctgca  
541 ccgcccagct tcccgggatg agggcccccg gtgtgggtcac ccggcgcgcc ccaggtcgct  
601 gagggaaccc ggccaggcgc ggagatgggg gtgcacggtg agtactcgcg ggtggggcgc  
661 tcccgcgccc cgggtccctg tttgagcggg gatttagcgc cccggctatt ggccaggagg  
721 tggctggggt caaggaccgg cgaactgtca aggaccccg aagggggagg ggggtggggc  
781 agcctccacg tgcagcggg gacttggggg agtccctggg gatggcaaaa acctgacctg  
841 tgaaggggac acagtttggg ggttggggg aagaaggttt gggggttctg ctgtgccagt  
901 ggagaggaag ctgataagct gataacctgg gcgctggag caccacttat ctgccagagg  
961 ggaagcctct gtcacaccag gattgaagtt tggccggaga agtggatgct ggtagctggg  
1021 ggtggggtgt gcacacggca gcaggattga atgaaggcca gggaggcagc acctgagtgc  
1081 ttgcatgggt ggggacagga aggacgagct ggggagagga cgtggggatg aaggaaagctg  
1141 tccctccaca gccacccttc tccctccccc cctgactctc agcctggcta tctgttctag  
1201 aatgtcctgc ctggctgtgg cttctcctgt cccctgctgc gctccctctg ggccctccag  
1261 tctggggcgc cccaccacgc ctcatctgtg acagccgagt cctggagagg tacctcttgg  
1321 agggcaagga ggccgagaat atcacggtga gaccccttcc ccagcacatt ccacagaact  
1381 cagctcagg gcttcaggga actcctccca gatccaggaa cctggcactt ggtttggggg  
1441 ggagttggga agctagacac tgcccccta cataagaata agtctggtgg ccccaaacca  
1501 tacctggaaa ctaggcaagg agcaaaagcca gcagatccta cggcctgtgg ggcagggcca  
1561 gagccttcag ggacccttga ctccccgggc tgtgtgcatt tcagacgggc tgtgctgaac  
1621 actgcagctt gaatgagaat atcactgtcc cagacaccaa agttaatttc tatgcctgga  
1681 agaggatgga ggtgagttcc tttttttttt tttttcctt cttttggaga atctcatttg  
1741 cgagcctgat tttggatgaa agggagaatg atcgggggaa aggtaaaatg gagcagcaga  
1801 gatgaggctg cctgggcgca gaggtcacg tctataatcc caggctgaga tggccgagat  
1861 gggagaattg cttgagccct ggagtttcag accaacctag gcagcatagt gagatcccc  
1921 atctctacaa acatttaaaa aaattagtc ggtgaagtgg tgcatgggtg tagtcccaga  
1981 tatttggaa gctgaggcgg gaggatcgct tgagcccagg aatttgaggc tgcagtgagc  
2041 tgtgatcaca ccactgcact ccagcctcag tgacagagtg aggcctgtgc tcaaaaaaga  
2101 aaagaaaaaa gaaaaataat gagggctgta tgggaatacat tcattattca ttcactcact  
2161 cactcactca ttcattcatt cattcattca acaagtctta ttgcatacct tctgtttgct  
2221 cagcttgggt cttggggctg ctgaggggca ggagggagag ggtgacatgg gtcagctgac  
2281 tccagagtc cactccctgt aggtcgggca gcaggccgta gaagcttggc agggcctggc  
2341 cctgctgtcg gaagctgtcc tgcggggcca gcccctgttg gtcaactctt cccagccgtg  
2401 ggagcccttg cagctgcatg tggataaagc cgtcagtggt cttcgcagcc tcaccactct  
2461 gcttcggggt ctgggagccc aggtgagtag gagcggacac ttctgcttgc ctttctgtg  
2521 agaaggggag aagggtctt ctaaggagta caggaactgt ccgtattcct tccctttctg  
2581 tggcagtgca gcgacctct gttttctcct tggcagaagg aagccatctc cctccagat  
2641 gcggcctcag ctgctccact ccgaacaatc actgctgaca ctttccgcaa actcttccga  
2701 gtctactcca atttctccg gggaaagctg aagctgtaca caggggaggc ctgcaggaca  
2761 ggggacagat gaccaggtgt gtccacctgg gcatatccac cactccctc accaacattg  
2821 cttgtgccac accctccccc gccactcctg aaccocgtcg agggcctctc agctcagcgc  
2881 cagcctgtcc catggacact ccagtgccag caatgacatc tcaggggcca gaggaactgt  
2941 ccagagagca actctgagat ctaaggatgt cacagggcca acttgagggc ccagagcagg  
3001 aagcattcag agagcagctt taaactcagg gacagagcca tgctgggaag acgcctgagc  
3061 tcaactggca ccctgcaaaa tttgatgcca ggacacgctt tggaggcgat ttacctgttt  
3121 tcgacgctac catcagggac aggatgacct ggagaactta ggtggcaagc tgtgacttct  
3181 ccaggtctca cgggcatggg cactcccttg gtggcaagag ccccttgac accggggtgg  
3241 tgggaaccat gaagacagga tgggggctgg cctctggctc tcatggggtc caagttttgt  
3301 gtattcttca acctcattga caagaactga aaccaccaat atgactcttg gcttttctgt  
3361 tttctgggaa cctccaaatc cctcggctct gtcccactcc tggcagcagt ggcagcgtc  
3421 caggtccggg aaatgagggg tggagggggc tgggcccctac gtgctgtctc acacagcctg  
3481 tctgacctct cgacctaccg gcctaggcca caagctctgc ctacgctggg caataaggtg  
3541 tctccattca aggcctcacc gcagtaaggc agctgccaac cctgcccagg gcaaggctgc  
3601 ag

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**FIGURE 3B**

MGVHECPAWLWLLLSLLSLPLGLPVLGAPPRLICDSRVLERYLLEAKEAENITTGCAEHC  
SLNENITVPDTKVNIFYAWKRMEVGQQAVEVWQGLALLSEAVLRGQALLVNSSQPWEPLQL  
HVDKAVSGLRSLTTLLRALGAQKEAISPPDAASAAPLRTITADTFRKLFRVYSNFLRGKL  
KLYTGEACRTGDR

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## FIGURE 4

AGCTTGCCCCG GGATGAGGGC CACCGGTGTG GTCACCCGGC GCGCCCCAGG TCGCTGAGGG 60  
ACCCCGGCCA GCGCGGAGA TGGGGGTGCA CGAATGTCCT GCCTGGCTGT GGCTTCTCCT 120  
GTCCCTGCTG TCGCTCCCTC TGGGCCTCCC AGTCCTGGGC GCCCCACCAC GCCTCATCTG 180  
TGACAGCCGA GTCCTGGAGA GGTACCTCTT GGAGGCCAAG GAGGCCGAGA ATATCACGAC 240  
GGGCTGTGCT GAACATTGCA GCTTGAATGA GAATATCACT GTCCCAGACA CCAAAGTTAA 300  
TTTCTATGCC TGAAGAGGA TGGAGGTCGG GCAGCAGGCC GTAGAAGTCT GGCAGGGCCT 360  
GGCCCTGCTG TCGGAAGCTG TCCTGCGGGG CCAGGCCCTG TTGGTCAACT CTTCCCAGCC 420  
GTGGGAGCCC CTGCAGCTGC ATGTGGATAA AGCCGTCAGT GGCCTTCGCA GCCTCACCAC 480  
TCTGCTTCGG GCTCTGGGAG CCCAGAAGGA AGCCATCTCC CCTCCAGATG CGGCCTCAGC 540  
TGCTCCACTC CGAACAATCA CTGCTGACAC TTTCCGAAA CTCTTCCGAG TCTACTCCAA 600  
TTTCCTCCGG GGAAAGCTGA AGCTGTACAC AGGGGAGGCC TGCAGGACAG GGGACAGATG 660  
ACCAGGTGTG TCCACCTGGG CATATCCACC ACCTCCCTCA CCAATATTGC TTGTGCCACA 720  
CCCTCCCCCG CCACTCCTGA ACCCCGTCGA GGGGCTCTCA GCTCAGCGCC AGCCTGTCCC 780  
ATGGACCA 788

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## FIGURE 5

1 AAAAAGATGA GGTAATTGTG TTTTATAAT TAAATATTTT ATAATTAAAA TATTTATAAT  
TTTTTCTACT CCATTAACAC AAAAATATTA ATTTATAAAA TATTAATTTT ATAAATATTA

61 TAAAATATTT ATAATTAAAT ATTTTATAAT TAAAATATTT ATAATTAAAT ATTTTATAAT  
ATTTTATAAA TATTAATTTA TAAAATATTA ATTTTATAAA TATTAATTTA TAAAATATTA

121 TAAAATATTT ATAATTAAAT ATTTTATAAT TAAAATATTT ATAATTAAAT ATTTTATAAT  
ATTTTATAAA TATTAATTTA TAAAATATTA ATTTTATAAA TATTAATTTA TAAAATATTA

181 TAAAATATTT ATAATTAAAT ATTTTATAAT TAAAATATTT ATAATTAAAT ATTTTATAAT  
ATTTTATAAA TATTAATTTA TAAAATATTA ATTTTATAAA TATTAATTTA TAAAATATTA

241 TAAAATATTT ATAATTAAAT ATTTTATAAT TAAAATGTTT ATAATTAAAT ATTTTATAAT  
ATTTTATAAA TATTAATTTA TAAAATATTA ATTTTACAAA TATTAATTTA TAAAATATTA

301 TAAAATGTTT ATAATTACAT ATTTTATAAT TAAAATGTTT ATAATTACAT ATTTTATAAT  
ATTTTACAAA TATTAATGTA TAAAATATTA ATTTTACAAA TATTAATGTA TAAAATATTA

361 TAAAATGTTT ATAATTACAT ATTTTATAAT TAAAATGTTT ATAATTACAT ATTTTATAAT  
ATTTTACAAA TATTAATGTA TAAAATATTA ATTTTACAAA TATTAATGTA TAAAATATTA

421 TAAAATGTTT ATAATTACAT ATTTTATAAT TACATATTTT ATAAAGTATT TATAATTACA  
ATTTTACAAA TATTAATGTA TAAAATATTA ATGTATAAAA TATTCATAA ATATTAATGT

481 TATTTTATAA TTAAAGTATT TATAATTACA TATTTTATAA TTAAAGTATT TATAATTACA  
ATAAAATATT AATTCATAA ATATTAATGT ATAAATATT AATTCATAA ATATTAATGT

541 TATTTTATAA TTCAATATTT TATAAATAGT TAAAAAGACG AGGAAAAAAT TAAAAAGACG  
ATAAAATATT AAGTTATAAA ATATTTATCA ATTTTCTGC TCCTTTTATA ATTTTCTGC

601 AGGTTATTGA TCTCAGGAAT TGTATTTGCC AAGTGAGAAG GAAAAAATAT TCACAAAGGC  
TCCAATAACT AGAGTCCTTA ACATAAACGG TTCACTCTC CTTTTTTATA AGTGTTTCCG

661 TTGTA  
AACAT

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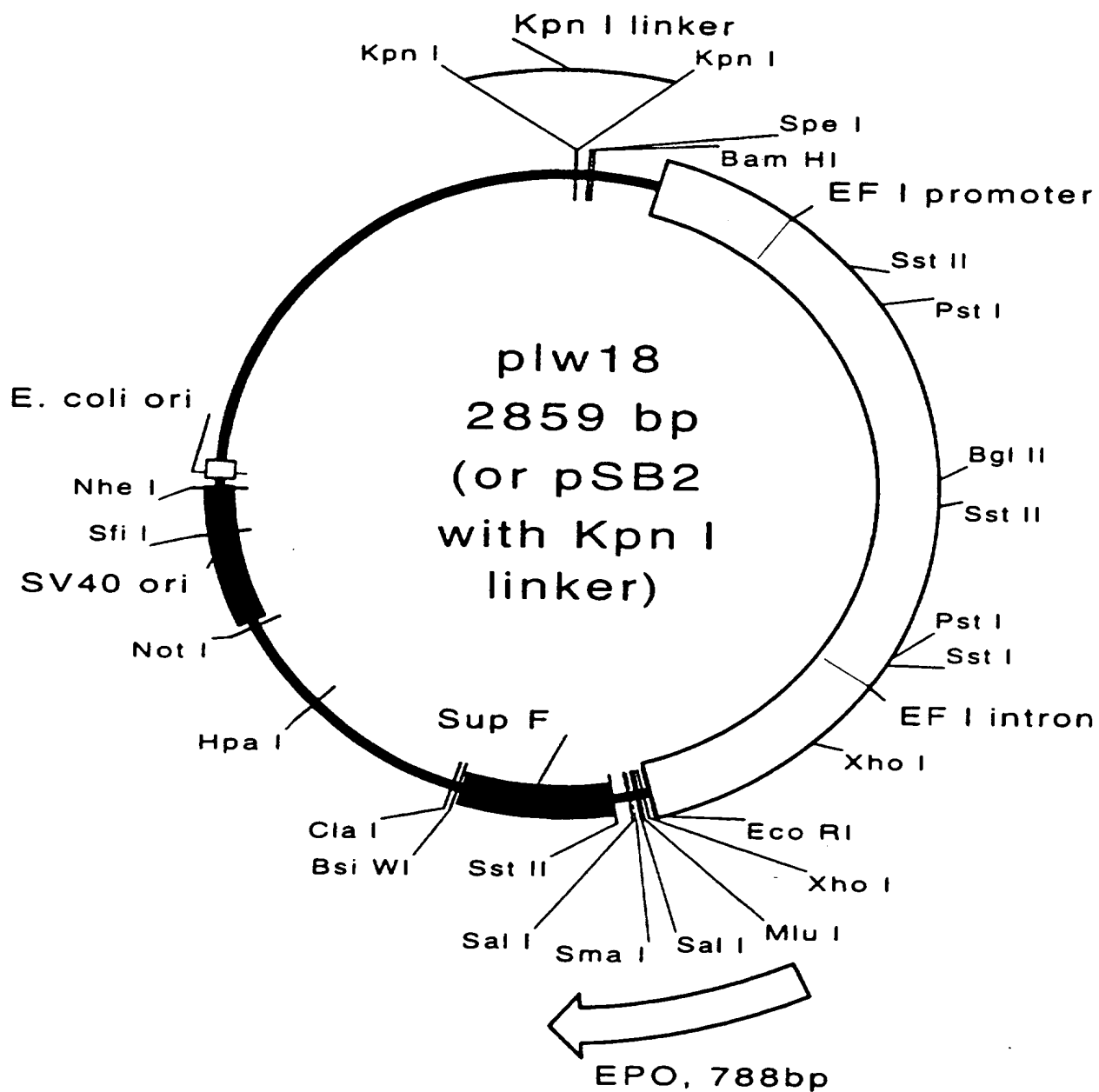
## FIGURE 6

1	TCTAGACCCC	AGTTTCTCTA	TAAGATGAGA	ATATTAGTCA	CGATTTGGTT	TCTAAGATCC
61	TGTCTATGTT	TGAGACTACA	GATACCTGTT	GCTACATTTT	CCTTCATAGC	TCTGAACAAG
121	GAGAATTCAG	CCCAATTCTC	ATGGCCTTCT	AAACAATCCA	GAGTTTCAGT	GCCATAAGGT
181	ACTACAATTT	AGTGTCAAAT	TAAAGTCAAAG	GCTTCATTAG	CCTGAAAGCT	CTGTCCCTGG
241	CCTGGGCATG	GCAAACGTGA	TCCCCCACTG	ACCATCCCCC	TGTCTCCCTT	CTCCCCAGAG
301	ACTCCAGTAG	CCTGGCGTCA	TCACAGGGGC	CAGACATATC	CAACATGTTT	CCAGCTTCCT
361	GCCACTTGAC	TTTCAGTGTG	CCTCCCTCTT	CAGTTACCCA	AATCCTGCCC	ACCATTCCAG
421	AGCCAGTTCA	ATCTCACCCA	TCCAGGACCC	CCGAGACCCC	CATCGTACCA	CTATAGTCTA
481	ACTGTGGTGT	AGACCCACA	CTGGGCACAT	TGCGTACGCT	CATTATTGGC	TGTGACGTCT
541	GATTATGCCC	TTCTCCTGGT	CTGGAAGCTC	TCGGAGGTGC	TCCATAATAC	ATGAAGAGAA
601	GTAGTGCTGG	TGTGGGAATA	GTGAGGTGTG	TTTATCCATC	CAGCTATCCG	GCACCAGCAC
661	TGGTCTCAGC	TTTCTGAGGT	AACACGTTCT	GAGCCTTAGT	CCTTGAGAGAA	CATAAAGAAA
721	ACTTTTTTTA	AAAGTAGTAA	AAAGTGGCTG	ACAAAAGCTG	ACCAAAAGCC	TTCAAAAGAA
781	ATGCTAAGTT	ATATCTAAGA	AAGTTTACCC	AAGGTCAGGC	AAATATGAAA	CCTAAAGCTA
841	GACGTGGGGA	AGAACTTCCG	GAGAGTTGCA	ATTCCCTGTG	CCCCAGCATC	CCCAGGAGGG
901	CATGCCCCACA	TCTGATTTAG	AAATCTGTGT	AAAATGAGTG	AAGGTTTCTA	TTTCTTGGGC
961	AGTGTGGGCA	CAGGTCTTTG	GAGAGGTCGA	TGGCCTCCCA	TAAAATCCTT	CCTGCTTGAT
1021	GGTTCTGGAT	CCTCAGCCAC	AGCTCCTAAT	AGCCATGAGG	TTTGAGCCCA	AAATAATTTA
1081	TGTGTTTGTT	TTTTCAGCCC	CAAAATTTCC	ATAGAATCAA	AGTAGTCAGA	GCTGAATGGG
1141	GCTAAGAGAC	CGTCCATTCC	TGCTTCTCA	TCACAGATGA	GGGACTGCCA	CCCAGAGCCC
1201	TAGAACTGT	CCCATGGCCC	CAGTTCCCAG	ACCCCTCCCT	TCTCCTACAG	CTCCAAGTTC
1261	ACTGTGCATT	CTAAATGAAG	ATGTAAACAT	AGGCAGCAAC	ACTCAAGAGT	AAAAATGAAG
1321	TGTGCATATG	AAAGAAAACCT	ATTCACATGG	ACCATATTAC	ATTATAATCA	CAGTGTTTAC
1381	TGCTTGACTA	CCATCTGCCT	GGGCTAGCAA	GGGTGTCAGT	GAGGAAGAGA	GGACAAGGGG
1441	TACCAATCTG	TGAACATACAC	ATGGTTCTTG	CTCTCCCAGC	TTCTCTCTCC	CATTGGCAAG
1501	GCAACAGGTA	AACACATGAA	AAATCAAATA	ATGCTATAAG	AGAAAAATGT	ATTCAGGACA
1561	ACAACAGGTT	TGTATGAAGG	CCTTTTCATCA	TCGTTGTCCT	ACCTAGAAAC	TGAATGACAG
1621	GGAATCAGAG	TCACAAGCTA	TGAAGTCTAA	CTGGGCTGGT	CCCAGAGAAA	GATTTCAGTGC
1681	AGTAGGTGGG	GCTGCAGCCA	GCCCTGGGTG	GGTGGAAGGA	TGACATCCAC	ATAGGCAAGA
1741	GGGTGATAAT	TCACTTGCGC	AGCTCCTCAC	TGCACATTGA	ACCCTGCTGA	CTTCTGGCTT
1801	CTCTCCCGGG	AGGAACTGCG	ACTCAACATT	CTGACCTTAT	CTCTTGGGTA	GCAGAATGAT
1861	GGAGAAGGAA	AGTTTCTTTT	TGCTTCTCGC	AGGGGTAAAT	CATCCATCTG	GAATGCCTAC
1921	ATTTGGTTGA	CAATGGCTCA	CCCTATCATC	TTCCTCCTGA	ACCATTACAC	TAAATGTGCC
1981	ATTTCTTTCC	TGATAGTTCT	CATTTGTGTG	TGTGTGTGTG	TGTGTGTGTG	TGCACGTGCT
2041	CACACATGCA	TGCTGTCACT	GGGTAAACAG	GCCACCCTGG	GCACAGTTCC	ATCTACAATG
2101	TTTGAAGTTT	ACTTTCCAGC	TTCTGGGCAT	CATTTGCAAT	TATAATGCTG	TCACAGGCAG
2161	AAACGAGATA	GGCTAATTAA	TCGTTGTCAA	TACTGATCCC	TATTTGCCAG	ATGAGATTTT
2221	GGAGCAGCAT	GGCTGGGAAT	AATTGGTATA	GACTGTATTT	CCTTGCTTTA	TGTCACTGGA
2281	AATATTTATT	TAAGCATCAC	GGTCGTATG	CATAAATATC	CTGGAAAATG	GGGTATAGCT
2341	GAATGGTGCA	GATTCAATTCA	TTCAATTATCA	GCAAATTATG	TTCTAAGCAC	CTACTTCAGT
2401	ACGTGAACAG	CACTAAACTC	AGAATATTGG	TCTGCTGGGG	TCCTTTATTA	GCTTCCATGA
2461	TTCCCTGAAC	TTGGCCAAGA	CCCTTCTGGT	CGGCTGCAGA	TAGGCACAAT	GGATAGTTTT
2521	GCTTCTAGA					

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FIGURE 7

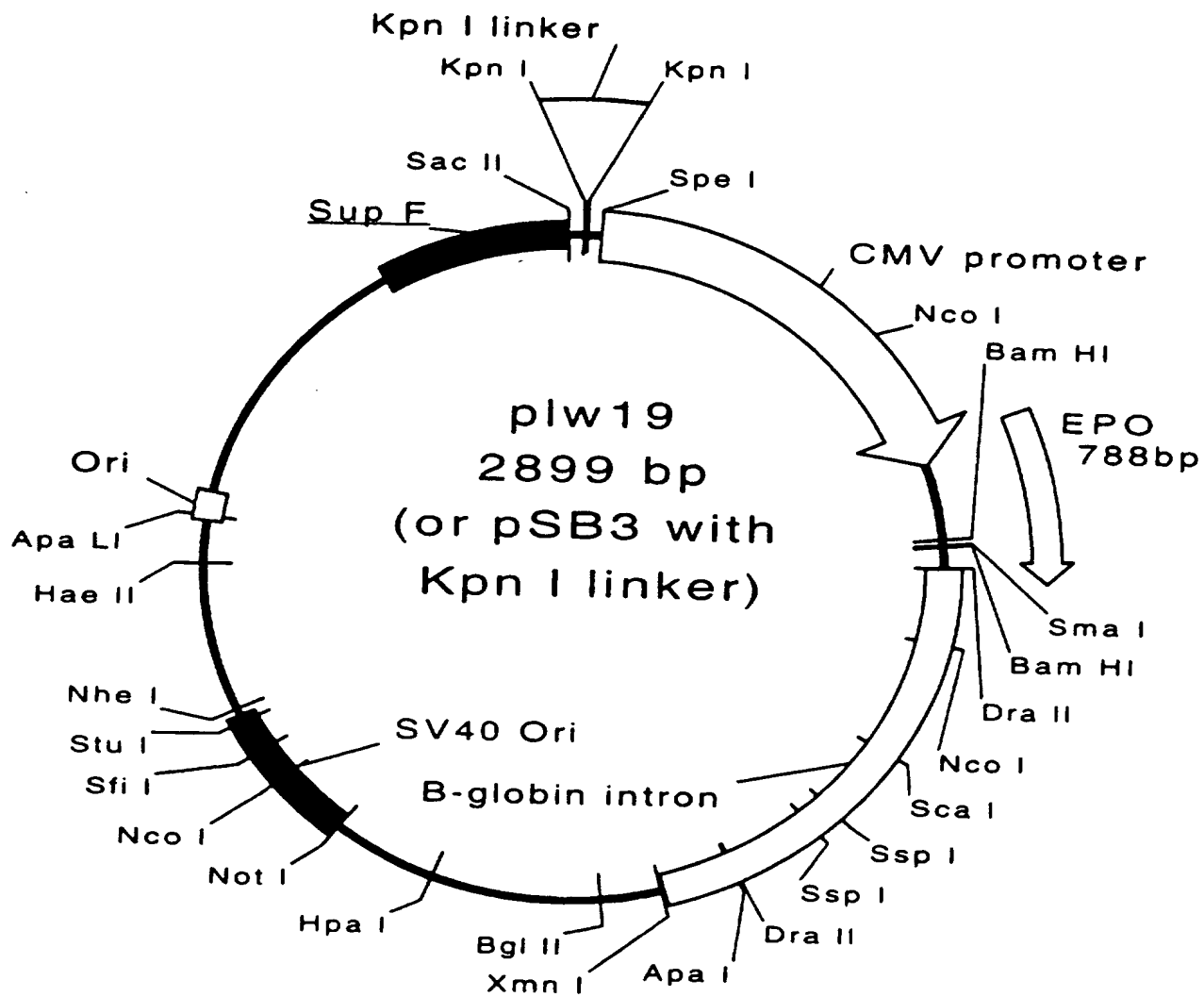


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FIGURE 8



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## INTERNATIONAL SEARCH REPORT

International Application No  
CA 95/00696

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C12N15/12 C12N15/67 C12N15/85 C12N5/10 C07K14/505  
A01K67/027

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C12N C07K A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	ANN. N.Y. ACAD. SCI., VOL. 665 (BIOCHEM. ENG. VII), 1992 pages 127-136, E.I. TSAO ET AL. 'Optimization of a roller bottle precess for the production of recombinant erythropoietin' see the whole document ---	20
X	PROC. NATL. ACAD. SCI., vol. 82, November 1985 NATL. ACAD SCI., WASHINGTON, DC, US;, pages 7580-7584, F.-K. LIN ET AL. 'Cloning and expression of human erythropoietin gene' see the whole document ---	20
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☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

## \* Special categories of cited documents:

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Date of the actual completion of the international search

29 March 1996

Date of mailing of the international search report

02.04.96

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## INTERNATIONAL SEARCH REPORT

International Application No.  
PCT/CA 95/00696

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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Information on patent family members

International Application No

PCT/CA 95/00696

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